

GenCore version 5.1.6  
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OM nucleic - nucleic search, using sw model  
Run on: August 19, 2004, 03:11:30 ; Search time 7823 Seconds  
(without alignments)  
11418.892 Million cell updates/sec  
Title: US-09-930-591-1  
Perfect score: 2061  
Sequence: 1 atggcgctatcacggccta.....atgaaatggaagagtgtga 2061

Scoring table: OLIGO NUC  
Gapop 60.0 , Gapext 60.0  
Searched: 3470272 seqs, 21671516995 residues  
Word size : 35 *or more*  
Total number of hits satisfying chosen parameters: 311

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000  
Post-processing: Listing first 1000 summaries  
Database : GenEmbl.\*  
1: gb.ba.\*  
2: gb.htg.\*  
3: gb.in.\*  
4: gb.om.\*  
5: gb.ov.\*  
6: gb.pat.\*  
7: gb.ph.\*  
8: gb.pl.\*  
9: gb.pr.\*  
10: gb.ro.\*  
11: gb.sts.\*  
12: gb.sy.\*  
13: gb.un.\*  
14: gb.vi.\*  
15: em.ba.\*  
16: em.fun.\*  
17: em.hum.\*  
18: em.in.\*  
19: em.mu.\*  
20: em.om.\*  
21: em.or.\*  
22: em.ov.\*  
23: em.pat.\*  
24: em.ph.\*  
25: em.pl.\*  
26: em.ro.\*  
27: em.sts.\*  
28: em.un.\*  
29: em.vi.\*  
30: em.htg.hum.\*  
31: em.htg.inv.\*  
32: em.htg.other.\*  
33: em.htg.mus.\*  
34: em.htg.pln.\*  
35: em.htg.rod.\*  
36: em.htg.man.\*  
37: em.htg.vrt.\*  
38: em.sy.\*  
39: em.htgo.hum.\*  
40: em.htgo.mus.\*  
41: em.htgo.other.\*

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB	ID	Description
1	2061	100.0	2061	6	AX441176	AX441176 Sequence
2	2061	100.0	2061	6	AX467113	AX467113 Sequence
3	92	4.5	9610	14	HEC278830	AJ78830 Hepatitis
4	76	3.7	9426	14	AF511949	AF511949 Hepatitis
5	62	3.0	943	6	A22779	A22779 non-structu
6	62	3.0	943	6	AR031209	AR031209 Sequence
7	62	3.0	943	6	AR145025	AR145025 Sequence
8	62	3.0	943	6	AR145025	AR145025 Sequence
9	62	3.0	9365	14	HPCNS3NS4	D10664 Hepatitis C
10	62	3.0	9365	14	AF290978	AF290978 Hepatitis
11	61	3.0	9395	14	AF511950	AF511950 Hepatitis
12	59	2.9	241	14	BD226202	BD226202 Improved
13	59	2.9	241	14	HPCNS10CLN	M94401 Hepatitis C
14	59	2.9	241	14	HPCNS11CLN	M94401 Hepatitis C
15	59	2.9	550	14	HCVNSTP	X71406 Hepatitis C
16	58	2.8	1477	14	HPCNS34	M60220 Hepatitis C
17	58	2.8	382	6	AR124773	AR124773 Sequence
18	58	2.8	382	6	AR353560	AR353560 Sequence
19	58	2.8	1414	6	AR124772	AR124772 Sequence
20	58	2.8	1414	6	AR353559	AR353559 Sequence
21	58	2.8	1420	6	AR124774	AR124774 Sequence
22	57	2.8	1420	6	AR353561	AR353561 Sequence
23	57	2.8	162	14	HPCNS35	M55151 Hepatitis C
24	57	2.8	223	14	HPCNS17CLN	M94451 Hepatitis C
25	57	2.8	229	14	HPCNS15CLN	M94449 Hepatitis C
26	56	2.7	281	6	HPCNSCLN5	M94469 Hepatitis C
27	56	2.7	281	6	I32190	I32190 Sequence 75
28	56	2.7	281	6	I34281	I34281 Sequence 75
29	56	2.7	281	6	I82486	I82486 Sequence 75
30	56	2.7	281	6	BD140411	BD140411 Hepatitis
31	56	2.7	368	6	I32188	AR118676 Sequence
32	56	2.7	368	6	I34279	I34279 Sequence 71
33	56	2.7	368	6	I82484	I82484 Sequence 71
34	56	2.7	368	6	BD140409	BD140409 Hepatitis
35	56	2.7	543	14	AF369231	AF369231 Hepatitis
36	56	2.7	1284	14	HPCNS3PROB	M62386 Hepatitis C
37	56	2.7	2058	6	AR404933	AR404933 Sequence
38	56	2.7	2058	6	AR408362	AR408362 Sequence
39	56	2.7	2058	6	AX395309	AX395309 Sequence
40	56	2.7	2058	6	AX454818	AX454818 Sequence
41	56	2.7	2064	6	I32187	I32187 Sequence 69
42	56	2.7	2064	6	I34278	I34278 Sequence 69
43	56	2.7	2064	6	I82483	I82483 Sequence 69
44	56	2.7	2064	6	BD140408	BD140408 Hepatitis
45	56	2.7	2283	6	BD140416	BD140416 Hepatitis
46	56	2.7	2523	6	I32195	I32195 Sequence 85
47	56	2.7	2523	6	I34286	I34286 Sequence 85
48	56	2.7	2523	6	I82491	I82491 Sequence 85
49	56	2.7	5360	6	AR118666	AR118666 Sequence
50	56	2.7	5360	6	I06434	I06434 Sequence 48
51	56	2.7	5360	6	I09328	I09328 Sequence 8
52	56	2.7	6299	6	AX164584	AX164584 Sequence
53	56	2.7	6785	6	AR118692	AR118692 Sequence
54	56	2.7	6785	6	I06440	I06440 Sequence 54
55	56	2.7	6785	6	I09329	I09329 Sequence 10
56	56	2.7	7310	6	AR118696	AR118696 Sequence
57	56	2.7	7310	6	I09331	I09331 Sequence 15
58	56	2.7	7310	14	HPCPOLYP	M32084 Hepatitis C
59	56	2.7	8316	6	AR118703	AR118703 Sequence
60	56	2.7	8987	6	AR118728	AR118728 Sequence
61	56	2.7	9185	6	AR118722	AR118722 Sequence
62	56	2.7	9185	6	AR118723	AR118723 Sequence
63	56	2.7	9185	6	I08294	I08294 Sequence 1
64	56	2.7	9185	6	BD091382	BD091382 HCV culti
65	56	2.7	9379	6	AR118747	AR118747 Sequence

Pred. No. is the number of results predicted by chance to have a

66	56	2.7	9379	6	AR166930	Sequence	139	48	2.3	6935	14	HPCT2	D16435	Hepatitis C
67	56	2.7	9379	6	AR301300	Sequence	140	48	2.3	9033	14	D89872	D89872	Hepatitis C
68	56	2.7	9401	6	AR176483	Sequence	141	48	2.3	9379	14	AF207765	AF207765	Hepatitis C
69	56	2.7	9401	6	B66593	Hepatitis C	142	48	2.3	9427	14	HPCURNA	D14484	Hepatitis C
70	56	2.7	9401	6	B71894	Sequence 9	143	48	2.3	9434	14	HPCUTB	D11355	Hepatitis C
71	56	2.7	9401	6	B11885	Sequence 9	144	48	2.3	9436	6	E07266	E07266	Blood-spleen
72	56	2.7	9401	6	BD080334	Hepatitis C	145	48	2.3	9436	14	HPCUTB	D11168	Hepatitis C
73	56	2.7	9401	6	AF378780	Synthetic	146	47	2.3	232	14	HPCNS7CLN	M94411	Hepatitis C
74	56	2.7	9609	12	AF378780	Synthetic	147	47	2.3	241	14	HPCNS4CLN	M94402	Hepatitis C
75	56	2.7	9609	12	AF378780	Synthetic	148	47	2.3	543	14	AF369232	AF369232	Hepatitis C
76	56	2.7	9609	12	AF378780	Synthetic	149	47	2.3	543	14	AF369242	AF369242	Hepatitis C
77	56	2.7	9693	12	AF378780	Synthetic	150	47	2.3	543	14	AF369243	AF369243	Hepatitis C
78	55	2.7	9693	12	AF378780	Synthetic	151	47	2.3	543	14	AF369243	AF369243	Hepatitis C
79	53	2.6	414	14	HCUI4261	Sequence	152	47	2.3	9502	6	E08263	E08263	GRNA of Hep
80	53	2.6	475	6	AX361030	Sequence	153	47	2.3	9502	6	E08264	E08264	cDNA of Hep
81	53	2.6	475	6	AX361032	Sequence	154	47	2.3	9502	6	E08264	E08264	cDNA of Hep
82	53	2.6	583	6	AX377698	Sequence	155	44	2.1	9502	6	E08264	E08264	cDNA of Hep
83	53	2.6	583	6	AX377698	Sequence	156	44	2.1	9502	6	E08264	E08264	cDNA of Hep
84	53	2.6	585	14	S68681	putative no	157	44	2.1	9502	6	E08264	E08264	cDNA of Hep
85	53	2.6	790	6	AX361031	Sequence	158	44	2.1	9502	6	E08264	E08264	cDNA of Hep
86	53	2.6	790	6	AX377697	Sequence	159	44	2.1	9502	6	E08264	E08264	cDNA of Hep
87	53	2.6	836	6	AX361029	Sequence	160	44	2.1	9502	6	E08264	E08264	cDNA of Hep
88	53	2.6	836	6	AX377695	Sequence	161	44	2.1	9502	6	E08264	E08264	cDNA of Hep
89	53	2.6	1284	14	HPCNS3PROA	Sequence	162	44	2.1	9502	6	E08264	E08264	cDNA of Hep
90	53	2.6	9416	6	AX441173	Sequence	163	44	2.1	9502	6	E08264	E08264	cDNA of Hep
91	53	2.6	9416	6	AX441173	Sequence	164	44	2.1	9502	6	E08264	E08264	cDNA of Hep
92	53	2.6	9518	6	AX100563	Sequence	165	44	2.1	9502	6	E08264	E08264	cDNA of Hep
93	53	2.6	9599	6	AR119831	Sequence	166	44	2.1	9502	6	E08264	E08264	cDNA of Hep
94	53	2.6	9599	6	AR119831	Sequence	167	44	2.1	9502	6	E08264	E08264	cDNA of Hep
95	53	2.6	9599	6	AR119831	Sequence	168	44	2.1	9502	6	E08264	E08264	cDNA of Hep
96	53	2.6	9599	6	AR119831	Sequence	169	44	2.1	9502	6	E08264	E08264	cDNA of Hep
97	53	2.6	9599	6	AR119831	Sequence	170	44	2.1	9502	6	E08264	E08264	cDNA of Hep
98	53	2.6	9599	6	AR119831	Sequence	171	44	2.1	9502	6	E08264	E08264	cDNA of Hep
99	53	2.6	9599	6	AR119831	Sequence	172	44	2.1	9502	6	E08264	E08264	cDNA of Hep
100	53	2.6	9599	6	AR119831	Sequence	173	44	2.1	9502	6	E08264	E08264	cDNA of Hep
101	53	2.6	9599	6	AR119831	Sequence	174	44	2.1	9502	6	E08264	E08264	cDNA of Hep
102	53	2.6	9599	6	AR119831	Sequence	175	44	2.1	9502	6	E08264	E08264	cDNA of Hep
103	53	2.6	9599	6	AR119831	Sequence	176	44	2.1	9502	6	E08264	E08264	cDNA of Hep
104	53	2.6	9599	6	AR119831	Sequence	177	44	2.1	9502	6	E08264	E08264	cDNA of Hep
105	53	2.6	9599	6	AR119831	Sequence	178	44	2.1	9502	6	E08264	E08264	cDNA of Hep
106	53	2.6	9599	6	AR119831	Sequence	179	44	2.1	9502	6	E08264	E08264	cDNA of Hep
107	53	2.6	9599	6	AR119831	Sequence	180	44	2.1	9502	6	E08264	E08264	cDNA of Hep
108	53	2.6	9599	6	AR119831	Sequence	181	44	2.1	9502	6	E08264	E08264	cDNA of Hep
109	53	2.6	9599	6	AR119831	Sequence	182	44	2.1	9502	6	E08264	E08264	cDNA of Hep
110	53	2.6	9599	6	AR119831	Sequence	183	44	2.1	9502	6	E08264	E08264	cDNA of Hep
111	52	2.5	885	6	AX48835	Sequence 1	184	44	2.1	9502	6	E08264	E08264	cDNA of Hep
112	52	2.5	885	6	AX48835	Sequence 1	185	44	2.1	9502	6	E08264	E08264	cDNA of Hep
113	52	2.5	885	6	AX48835	Sequence 1	186	44	2.1	9502	6	E08264	E08264	cDNA of Hep
114	52	2.5	885	6	AX48835	Sequence 1	187	44	2.1	9502	6	E08264	E08264	cDNA of Hep
115	52	2.5	885	6	AX48835	Sequence 1	188	44	2.1	9502	6	E08264	E08264	cDNA of Hep
116	52	2.5	885	6	AX48835	Sequence 1	189	44	2.1	9502	6	E08264	E08264	cDNA of Hep
117	50	2.4	337	6	AX360976	Sequence	190	44	2.1	9502	6	E08264	E08264	cDNA of Hep
118	50	2.4	337	6	AX360976	Sequence	191	44	2.1	9502	6	E08264	E08264	cDNA of Hep
119	50	2.4	337	6	AX360976	Sequence	192	44	2.1	9502	6	E08264	E08264	cDNA of Hep
120	50	2.4	337	6	AX360976	Sequence	193	44	2.1	9502	6	E08264	E08264	cDNA of Hep
121	50	2.4	337	6	AX360976	Sequence	194	44	2.1	9502	6	E08264	E08264	cDNA of Hep
122	50	2.4	337	6	AX360976	Sequence	195	44	2.1	9502	6	E08264	E08264	cDNA of Hep
123	50	2.4	337	6	AX360976	Sequence	196	44	2.1	9502	6	E08264	E08264	cDNA of Hep
124	50	2.4	337	6	AX360976	Sequence	197	44	2.1	9502	6	E08264	E08264	cDNA of Hep
125	50	2.4	337	6	AX360976	Sequence	198	44	2.1	9502	6	E08264	E08264	cDNA of Hep
126	50	2.4	337	6	AX360976	Sequence	199	44	2.1	9502	6	E08264	E08264	cDNA of Hep
127	50	2.4	337	6	AX360976	Sequence	200	44	2.1	9502	6	E08264	E08264	cDNA of Hep
128	50	2.4	337	6	AX360976	Sequence	201	44	2.1	9502	6	E08264	E08264	cDNA of Hep
129	50	2.4	337	6	AX360976	Sequence	202	44	2.1	9502	6	E08264	E08264	cDNA of Hep
130	50	2.4	337	6	AX360976	Sequence	203	44	2.1	9502	6	E08264	E08264	cDNA of Hep
131	50	2.4	337	6	AX360976	Sequence	204	44	2.1	9502	6	E08264	E08264	cDNA of Hep
132	50	2.4	337	6	AX360976	Sequence	205	44	2.1	9502	6	E08264	E08264	cDNA of Hep
133	50	2.4	337	6	AX360976	Sequence	206	44	2.1	9502	6	E08264	E08264	cDNA of Hep
134	50	2.4	337	6	AX360976	Sequence	207	44	2.1	9502	6	E08264	E08264	cDNA of Hep
135	50	2.4	337	6	AX360976	Sequence	208	44	2.1	9502	6	E08264	E08264	cDNA of Hep
136	50	2.4	337	6	AX360976	Sequence	209	44	2.1	9502	6	E08264	E08264	cDNA of Hep
137	48	2.3	5211	6	BD270487	Substitut	210	44	2.1	9502	6	E08264	E08264	cDNA of Hep
138	48	2.3	5211	6	AX044453	Sequence	211	44	2.1	9502	6	E08264	E08264	cDNA of Hep

212	44	2.1	19912	6	AX164586	285	1.7	162	14	HPCHCV49	M60338 Hepatitis C
213	44	2.1	20160	6	AX164590	286	1.7	287	6	AR037528	AR037528 Sequence
214	44	2.1	20217	6	AX164594	287	1.7	287	6	E09291	E09291 DNA encodin
215	44	2.1	20247	6	AX164596	288	1.7	372	14	AF379284	AF379284 Hepatitis
216	44	2.1	20316	6	AX164592	289	1.7	372	14	AF379284	AF379284 Sequence
217	43	2.1	347	6	AR118677	290	1.7	390	6	AX361036	AX361036 Sequence
218	43	2.1	347	6	AR118677	291	1.7	390	6	AX361036	AX361036 Sequence
219	43	2.1	2748	6	E09628	292	1.7	414	14	HCUI4260	U14260 Hepatitis C
220	43	2.1	3559	14	HPCPNS234S	293	1.7	414	14	HCUI4279	U14279 Hepatitis C
221	42	2.0	414	14	HCUI4270	294	1.7	453	6	AX360952	AX360952 Sequence
222	41	2.0	268	6	AR118670	295	1.7	453	6	AX360952	AX360952 Sequence
223	41	2.0	307	6	AR118670	296	1.7	475	6	AX361035	AX361035 Sequence
224	41	2.0	308	6	AR118733	297	1.7	475	6	AX361035	AX361035 Sequence
225	41	2.0	308	6	AR118733	298	1.7	495	6	AX361035	AX361035 Sequence
226	41	2.0	308	6	AR118733	299	1.7	495	6	AX361035	AX361035 Sequence
227	41	2.0	308	6	AR118733	300	1.7	520	14	HPCHCV49	HPCHCV49 Sequence
228	41	2.0	308	6	AR118733	301	1.7	520	14	HPCHCV49	HPCHCV49 Sequence
229	41	2.0	477	6	AR061073	302	1.7	520	14	HPCHCV49	HPCHCV49 Sequence
230	41	2.0	477	6	AR061073	303	1.7	520	14	HPCHCV49	HPCHCV49 Sequence
231	41	2.0	477	6	AR061073	304	1.7	520	14	HPCHCV49	HPCHCV49 Sequence
232	41	2.0	477	6	AR061073	305	1.7	520	14	HPCHCV49	HPCHCV49 Sequence
233	41	2.0	477	6	AR061073	306	1.7	520	14	HPCHCV49	HPCHCV49 Sequence
234	41	2.0	477	6	AR061073	307	1.7	520	14	HPCHCV49	HPCHCV49 Sequence
235	41	2.0	477	6	AR061073	308	1.7	520	14	HPCHCV49	HPCHCV49 Sequence
236	41	2.0	477	6	AR061073	309	1.7	520	14	HPCHCV49	HPCHCV49 Sequence
237	41	2.0	480	6	BD091383	310	1.7	9374	14	AF207753	AF207753 Hepatitis
238	41	2.0	495	6	BD091383	311	1.7	9374	14	AF207753	AF207753 Sequence
239	41	2.0	495	6	BD091383	312	1.7	9374	14	AF207753	AF207753 Sequence
240	41	2.0	495	6	BD091383	313	1.7	9374	14	AF207753	AF207753 Sequence
241	41	2.0	495	6	BD091383	314	1.7	9374	14	AF207753	AF207753 Sequence
242	41	2.0	495	6	BD091383	315	1.7	9374	14	AF207753	AF207753 Sequence
243	41	2.0	543	14	AF369239	316	1.7	9374	14	AF207753	AF207753 Sequence
244	41	2.0	543	14	AF369239	317	1.7	9374	14	AF207753	AF207753 Sequence
245	41	2.0	543	14	AF369239	318	1.7	9374	14	AF207753	AF207753 Sequence
246	41	2.0	543	14	AF369239	319	1.7	9374	14	AF207753	AF207753 Sequence
247	41	2.0	543	14	AF369239	320	1.7	9374	14	AF207753	AF207753 Sequence
248	41	2.0	543	14	AF369239	321	1.7	9374	14	AF207753	AF207753 Sequence
249	41	2.0	543	14	AF369239	322	1.7	9374	14	AF207753	AF207753 Sequence
250	41	2.0	543	14	AF369239	323	1.7	9374	14	AF207753	AF207753 Sequence
251	41	2.0	543	14	AF369239	324	1.7	9374	14	AF207753	AF207753 Sequence
252	41	2.0	543	14	AF369239	325	1.7	9374	14	AF207753	AF207753 Sequence
253	41	2.0	543	14	AF369239	326	1.7	9374	14	AF207753	AF207753 Sequence
254	41	2.0	543	14	AF369239	327	1.7	9374	14	AF207753	AF207753 Sequence
255	41	2.0	543	14	AF369239	328	1.7	9374	14	AF207753	AF207753 Sequence
256	41	2.0	543	14	AF369239	329	1.7	9374	14	AF207753	AF207753 Sequence
257	41	2.0	543	14	AF369239	330	1.7	9374	14	AF207753	AF207753 Sequence
258	41	2.0	543	14	AF369239	331	1.7	9374	14	AF207753	AF207753 Sequence
259	41	2.0	543	14	AF369239	332	1.7	9374	14	AF207753	AF207753 Sequence
260	41	2.0	543	14	AF369239	333	1.7	9374	14	AF207753	AF207753 Sequence
261	41	2.0	543	14	AF369239	334	1.7	9374	14	AF207753	AF207753 Sequence
262	41	2.0	543	14	AF369239	335	1.7	9374	14	AF207753	AF207753 Sequence
263	41	2.0	543	14	AF369239	336	1.7	9374	14	AF207753	AF207753 Sequence
264	41	2.0	543	14	AF369239	337	1.7	9374	14	AF207753	AF207753 Sequence
265	41	2.0	543	14	AF369239	338	1.7	9374	14	AF207753	AF207753 Sequence
266	41	2.0	543	14	AF369239	339	1.7	9374	14	AF207753	AF207753 Sequence
267	39	1.9	9538	14	HCUB9019	340	1.7	9538	14	HCUB9019	HCUB9019 Sequence
268	38	1.8	372	14	AF379308	341	1.7	9538	14	HCUB9019	HCUB9019 Sequence
269	38	1.8	543	14	AF369234	342	1.7	9538	14	HCUB9019	HCUB9019 Sequence
270	38	1.8	543	14	AF369234	343	1.7	9538	14	HCUB9019	HCUB9019 Sequence
271	38	1.8	9391	6	E04421	344	1.7	9538	14	HCUB9019	HCUB9019 Sequence
272	38	1.8	9391	6	E04421	345	1.7	9538	14	HCUB9019	HCUB9019 Sequence
273	38	1.8	9440	14	S62220	346	1.7	9538	14	HCUB9019	HCUB9019 Sequence
274	38	1.8	9440	14	S62220	347	1.7	9538	14	HCUB9019	HCUB9019 Sequence
275	38	1.8	9487	6	E08443	348	1.7	9538	14	HCUB9019	HCUB9019 Sequence
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284	35	1.7	543	14	HPCHCV36	357	1.7	9573	14	HPCCGS	HPCCGS Sequence
285	35	1.7	543	14	HPCHCV36	358	1.7	9573	14	HPCCGS	HPCCGS Sequence
286	35	1.7	543	14	HPCHCV36	359	1.7	9573	14	HPCCGS	HPCCGS Sequence
287	35	1.7	543	14	HPCHCV36	360	1.7	9573	14	HPCCGS	HPCCGS Sequence
288	35	1.7	543	14	HPCHCV36	361	1.7	9573	14	HPCCGS	HPCCGS Sequence
289	35	1.7	543	14	HPCHCV36	362	1.7	9573	14	HPCCGS	HPCCGS Sequence
290	35	1.7	543	14	HPCHCV36	363	1.7	9573	14	HPCCGS	HPCCGS Sequence
291	35	1.7	543	14	HPCHCV36	364	1.7	9573	14	HPCCGS	HPCCGS Sequence
292	35	1.7	543	14	HPCHCV36	365	1.7	9573	14	HPCCGS	HPCCGS Sequence
293	35	1.7	543	14	HPCHCV36	366	1.7	9573	14	HPCCGS	HPCCGS Sequence
294	35	1.7	543	14	HPCHCV36	367	1.7	9573	14	HPCCGS	HPCCGS Sequence
295	35	1.7	543	14	HPCHCV36	368	1.7	9573	14	HPCCGS	HPCCGS Sequence
296	35	1.7	543	14	HPCHCV36	369	1.7	9573	14	HPCCGS	HPCCGS Sequence
297	35	1.7	543	14	HPCHCV36	370	1.7	9573	14	HPCCGS	HPCCGS Sequence
298	35	1.7	543	14	HPCHCV36	371	1.7	9573	14	HPCCGS	HPCCGS Sequence
299	35	1.7	543	14	HPCHCV36	372	1.7	9573	14	HPCCGS	HPCCGS Sequence
300	35	1.7	543	14	HPCHCV36	373	1.7	9573	14	HPCCGS	HPCCGS Sequence
301	35	1.7	543	14	HPCHCV36	374	1.7	9573	14	HPCCGS	HPCCGS Sequence
302	35	1.7	543	14	HPCHCV36	375	1.7	9573	14	HPCCGS	HPCCGS Sequence
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304	35	1.7	543	14	HPCHCV36	377	1.7	9573	14	HPCCGS	HPCCGS Sequence
305	35	1.7	543	14	HPCHCV36	378	1.7	9573	14	HPCCGS	HPCCGS Sequence
306	35	1.7	543	14	HPCHCV36	379	1.7	9573	14	HPCCGS	HPCCGS Sequence
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323	35	1.7	543	14	HPCHCV36	396	1.7	9573	14	HPCCGS	HPCCGS Sequence
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325	35	1.7	543	14	HPCHCV36	398	1.7	9573	14	HPCCGS	HPCCGS Sequence
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327	35	1.7	543	14	HPCHCV36	400	1.7	9573	14	HP	

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## RESULT 2

AX467113

LOCUS

DEFINITION

Sequence 1 from Patent WO0214362.

ACCESSION

AX467113

VERSION

AX467113.1

KEYWORDS

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ORIGIN		/db_xref="taxon:32630" /note="Hepatitis C virus NS3/4A coding region"		Query Match Best Local Similarity 100.0%; Score 2061; DB 6; Length 2061; Matches 2061; Conservative 0; Mismatches 0; Indels 0; Gaps 0;	
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DB	1	ATGGCGCTATACCGGCTATGCCAGCAGACAAGGGGCTTTGGGATGCATTAATCACC	60		
QY	61	AGCTTGACCGCGCGGACAAAACACAGGTGAGGTGAGGTTCAGATCGTGCACATGCT	120		
DB	61	AGCTTGACCGCGCGGACAAAACACAGGTGAGGTGAGGTTCAGATCGTGCACATGCT	120		
QY	121	GCCAGACATTTCCTTGGCAACTCTGATTAACCGGGTGTGTTGGACTGTCTACCATGGAGCC	180		
DB	121	GCCAGACATTTCCTTGGCAACTCTGATTAACCGGGTGTGTTGGACTGTCTACCATGGAGCC	180		
QY	181	GGAAACAGGACCATTCGCTACCTAAGGTTCCTGTTATCCAGATGTACCAATGTGGAC	240		
DB	181	GGAAACAGGACCATTCGCTACCTAAGGTTCCTGTTATCCAGATGTACCAATGTGGAC	240		
QY	241	CAAGACCTCGTAGGCTGGCGGCTCCCAAGGTGCCCGCTCAITTAACACCATGCACTTGC	300		
DB	241	CAAGACCTCGTAGGCTGGCGGCTCCCAAGGTGCCCGCTCAITTAACACCATGCACTTGC	300		
QY	301	GGCTTCCTCGACCTTTACCTGGTCAAGGACACCGCGATGTCATTCCTGTGCGCCGACGG	360		
DB	301	GGCTTCCTCGACCTTTACCTGGTCAAGGACACCGCGATGTCATTCCTGTGCGCCGACGG	360		
QY	361	GCTGATGGCAGGGCAGCGCTTTGCGCCGCGCTATCTTACCTTGAAGGCTCCTCG	420		
DB	361	GCTGATGGCAGGGCAGCGCTTTGCGCCGCGCTATCTTACCTTGAAGGCTCCTCG	420		
QY	421	GGAGGCGCTCTGCTGTGCGCGCAGGACATCCCGTAGGCATATTCAGAGCGCGGTATGC	480		
DB	421	GGAGGCGCTCTGCTGTGCGCGCAGGACATCCCGTAGGCATATTCAGAGCGCGGTATGC	480		
QY	481	ACCGTGGAGTGGCTAAGCGGTGGACTTCATCCCGTAGAGCTTAGAGCAACCATG	540		
DB	481	ACCGTGGAGTGGCTAAGCGGTGGACTTCATCCCGTAGAGCTTAGAGCAACCATG	540		
QY	541	AGGTCCCGGTGTCTCAGACAATCTCTCCCAACAGCAGTGCCTCCAGAGCTACCAAGTG	600		
DB	541	AGGTCCCGGTGTCTCAGACAATCTCTCCCAACAGCAGTGCCTCCAGAGCTACCAAGTG	600		
QY	601	GCCACCTGATGCTCCACCGGAGCGGTAAAGACACCAAGTCCCGCGCATACGCA	660		
DB	601	GCCACCTGATGCTCCACCGGAGCGGTAAAGACACCAAGTCCCGCGCATACGCA	660		
QY	661	GCTCAGGGCTACAAAGTGTGCTCAACCCCTCGTTCGCTGCTCAACAATGGCTTTGGT	720		
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DB	721	GCTTACATGTCCAAAGGCCAATGGATGATCTTAACATCAGGACTGGGGTGAGGACAAT	780		
QY	781	ACTACTGGACCGGATCAGTATCCCTAGCGCAAGTTCCTTCCGACCGCGGTGT	840		
DB	781	ACTACTGGACCGGATCAGTATCCCTAGCGCAAGTTCCTTCCGACCGCGGTGT	840		
QY	841	TCAGGGGTGCTTATGACATAATAATTTGTGACAGTGCACCTCCACGGATGCAACATCC	900		
DB	841	TCAGGGGTGCTTATGACATAATAATTTGTGACAGTGCACCTCCACGGATGCAACATCC	900		
QY	901	ATCTTGGGCAATGGCACTGCTTGAACAGCAGACCGCGGGCGGAGACTGCTGTG	960		
DB	901	ATCTTGGGCAATGGCACTGCTTGAACAGCAGACCGCGGGCGGAGACTGCTGTG	960		
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DB	961	CTGCGCACCGCTACCCCTCCGGGCTCCGTCACTGTGCCCATCTCTAACATCGAGGAGTT	1020		
QY	1021	GCTCTGTCCACTACCGGAGAGATCCCTTTTATGGCAAGGCTATTCCTTGAAGCAAT	1080		
DB	1021	GCTCTGTCCACTACCGGAGAGATCCCTTTTATGGCAAGGCTATTCCTTGAAGCAAT	1080		
QY	1081	AAGGGGGGAGACATCTCATCTTCTGCCACTCAAAGAGAAGTGCAGAGCTCGCGCA	1140		
DB	1081	AAGGGGGGAGACATCTCATCTTCTGCCACTCAAAGAGAAGTGCAGAGCTCGCGCA	1140		
QY	1141	AAACTCGTCCGCTGGCGCTCAATGCCGTGCTTACTACCGCGGCTTATGTGTCCGTC	1200		
DB	1141	AAACTCGTCCGCTGGCGCTCAATGCCGTGCTTACTACCGCGGCTTATGTGTCCGTC	1200		
QY	1201	ATCCCGACAGTGGTGCAGCTGTGCTGCTGGCAACTGACGCGCTCATGACCGGCTTACC	1260		
DB	1201	ATCCCGACAGTGGTGCAGCTGTGCTGCTGGCAACTGACGCGCTCATGACCGGCTTACC	1260		
QY	1261	GGGCACTTCGANTCGGTGATAGACTGCAACAGTGTGTCAACAGACAGTGCAGTTCAGC	1320		
DB	1261	GGGCACTTCGANTCGGTGATAGACTGCAACAGTGTGTCAACAGACAGTGCAGTTCAGC	1320		
QY	1321	CTTGACCTTACCTTCAACATTCAGACATCAAGCTTCCCGAGGATGCTGTCCGCTACT	1380		
DB	1321	CTTGACCTTACCTTCAACATTCAGACATCAAGCTTCCCGAGGATGCTGTCCGCTACT	1380		
QY	1381	CAACGTCCGGGTAGGATGGCAGAGGGAAGCAGGATCTACAGATTTGTGCAACCGGG	1440		
DB	1381	CAACGTCCGGGTAGGATGGCAGAGGGAAGCAGGATCTACAGATTTGTGCAACCGGG	1440		
QY	1441	GAGCGTCTTCTGGCATGTTGACTCGTCTGTCTCTGCGAGTGTATGACCGGGTGT	1500		
DB	1441	GAGCGTCTTCTGGCATGTTGACTCGTCTGTCTCTGCGAGTGTATGACCGGGTGT	1500		
QY	1501	GCTTGTGTAGCTTACGCGCCCGAGACCAAGTTAGGCTACGAGCATACATGAACACC	1560		
DB	1501	GCTTGTGTAGCTTACGCGCCCGAGACCAAGTTAGGCTACGAGCATACATGAACACC	1560		
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DB	1561	CCGGACTTCCCGTGTGCCAAGACCATCTTGAATTTTGGAGGGCGTCTTTACGGGTCTC	1620		
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DB	1621	ACCCATAGAGCCGCTTCTCTATCCAGACAAAGCAGAGTGGGGAACCTTCCCTAT	1680		
QY	1681	CTGGTAGGTACCAAGCCACCGTGTGCGTAGAGCTCAAGCCCTTCCCGGCTCGTGGAC	1740		
DB	1681	CTGGTAGGTACCAAGCCACCGTGTGCGTAGAGCTCAAGCCCTTCCCGGCTCGTGGAC	1740		
QY	1741	CAGATGTGAAGTGTGTTGATCCGTCTCAAGCCCACTTCCATGGGCCAACCTCTGCTA	1800		
DB	1741	CAGATGTGAAGTGTGTTGATCCGTCTCAAGCCCACTTCCATGGGCCAACCTCTGCTA	1800		
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DB	1801	TATAGACTGGGCGTGTCCAGATGAAGTCAACCTGACGACCCAGTCAACAGTATATC	1860		
QY	1861	ATGACATGTATGTCCGCTGACCTGGAGGTCTGACAGATACCTGGGTCTCTGTTGGCGC	1920		
DB	1861	ATGACATGTATGTCCGCTGACCTGGAGGTCTGACAGATACCTGGGTCTCTGTTGGCGC	1920		
QY	1921	GTTCGGGTCTTTGGCGCGCTTATTCATCCAGGCTCGTGTGCTATAGTAGGTAGG	1980		
DB	1921	GTTCGGGTCTTTGGCGCGCTTATTCATCCAGGCTCGTGTGCTATAGTAGGTAGG	1980		
QY	1981	ATTGTCTTGTCCGAAAGCCGCAATCATACCCGACAGGGAAGTCTCTTACCGGAGTTC	2040		
DB	1981	ATTGTCTTGTCCGAAAGCCGCAATCATACCCGACAGGGAAGTCTCTTACCGGAGTTC	2040		
QY	2041	GATGAATGGAAGAGTGTCTGA	2061		

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# RESULT 3

HEC278830

LOCUS

DEFINITION

HEPATICITIS C VIRUS GENOMIC RNA FOR POLYPROTEIN GENE.

VERSION

HEC278830.1

KEYWORDS

core protein; envelop protein 1; envelop protein 2; non-structural protein 2; non-structural protein 3; non-structural protein 4a; non-structural protein 4b; non-structural protein 5a; non-structural protein 5b; ORF1; ORF2; ORF3; ORF4; ORF5; ORF6; ORF7; ORF8; ORF9; polyprotein.

## SOURCE

HEPATICITIS C VIRUS

ORGANISM

Viruses; serNA positive-strand viruses, no DNA stage; Flaviviridae; Hepacivirus.

## REFERENCE

1

Kumar, U., Tuthill, T., Thomas, H. C. and Morjardino, J.

Sequence, expression and reconstitution of an HCV genome from a

British isolate derived from a single blood donation

J. Viral Hepat. 7 (6), 459-465. (2000)

21014672

PUBMED

11115058

REFERENCE

2 (bases 1 to 9610)

Kumar, U.

AUTHORS

Direct Submission

Submitted (11-AUG-2000) Kumar U., Virology, GlaxoWellcome Research

centre, Gunneis Wood Road, Stevenage, Hertfordshire, SG1 2NY,

UNITED KINGDOM

LOCATION/Qualifiers

1. .9610

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/vixon

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342. .9377

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HEC278830 9610 bp RNA linear VRL 03-JAN-2001  
Hepatitis C virus genomic RNA for polyprotein gene.

AJ278830 GI:9843876  
core protein; envelop protein 1; envelop protein 2; non-structural protein 2; non-structural protein 3; non-structural protein 4a; non-structural protein 4b; non-structural protein 5a; non-structural protein 5b; ORF1; ORF2; ORF3; ORF4; ORF5; ORF6; ORF7; ORF8; ORF9; polyprotein.

Hepatitis C virus  
Viruses; serNA positive-strand viruses, no DNA stage; Flaviviridae; Hepacivirus.

1  
Kumar, U., Tuthill, T., Thomas, H. C. and Morjardino, J.  
Sequence, expression and reconstitution of an HCV genome from a  
British isolate derived from a single blood donation  
J. Viral Hepat. 7 (6), 459-465. (2000)

21014672  
PUBMED  
11115058

REFERENCE  
2 (bases 1 to 9610)  
Kumar, U.  
Direct Submission  
Submitted (11-AUG-2000) Kumar U., Virology, GlaxoWellcome Research  
centre, Gunneis Wood Road, Stevenage, Hertfordshire, SG1 2NY,  
UNITED KINGDOM

## FEATURES

source

## CDS

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QY 1579 CAAGACCATCTTGAAATTTGGAGGGGCTCTT 1610  
DB 4995 CAAGACCATCTTGAAATTTGGAGGGGCTCTT 5026

## RESULT 4

AF511949

LOCUS

DEFINITION

Hepatitis C virus isolate XF223 polyprotein gene, complete

sequence.

9426 bp RNA linear

VRL 13-JUN-2002

```

ACCESSION AF511949
VERSION AF511949.1 GI:21397076
KEYWORDS
SOURCE Hepatitis C virus
ORGANISM Hepatitis C virus
REFERENCE 1 (bases 1 to 9426)
AUTHORS Fan,X. and Di Bisceglie,A.M.
TITLE Clonal Nature of Hepatitis C Virus
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 9426)
AUTHORS Fan,X. and Di Bisceglie,A.M.
TITLE Direct Submission
JOURNAL Submitted (14-MAY-2002) Gastroenterology & Hepatology, Saint Louis
University School of Medicine, 1402 South Grand Blvd., St. Louis,
MO 63104, USA
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Db 4690 TCGGTGATAGACTGCAACACGTGTGTACCCAGACAGTCGACTTCAGCCCTTGACCCCTACC 4749
QY 1333 TTCACCATTTGAGACAA 1348
Db 4750 TTCACCATTTGAGACAA 4765

RESULT 5
LOCUS A22779 943 bp DNA linear PAT 24-JAN-1995
DEFINITION non-structural coding region.
ACCESSION A22779
VERSION A22779.1 GI:832940
KEYWORDS
SOURCE Hepatitis C virus
ORGANISM Hepatitis C virus
REFERENCE 1 (bases 1 to 943)
AUTHORS Brechot,C., Kreamsdorf,D. and Porchon,C.
TITLE Nucleotide and peptide sequences of an isolate of the hepatitis C
JOURNAL virus; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Hepacivirus.
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Db 630 GCGCGGCTTCTGGCTGTTGGCGCGGTATTGCGTATCCACAGGCTGCGTGCATAGTA 689
QY 1975 GG 1976

ACCESSION AF511949
VERSION AF511949.1 GI:21397076
KEYWORDS
SOURCE Hepatitis C virus
ORGANISM Hepatitis C virus
REFERENCE 1 (bases 1 to 9426)
AUTHORS Fan,X. and Di Bisceglie,A.M.
TITLE Clonal Nature of Hepatitis C Virus
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 9426)
AUTHORS Fan,X. and Di Bisceglie,A.M.
TITLE Direct Submission
JOURNAL Submitted (14-MAY-2002) Gastroenterology & Hepatology, Saint Louis
University School of Medicine, 1402 South Grand Blvd., St. Louis,
MO 63104, USA
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Db 4690 TCGGTGATAGACTGCAACACGTGTGTACCCAGACAGTCGACTTCAGCCCTTGACCCCTACC 4749
QY 1333 TTCACCATTTGAGACAA 1348
Db 4750 TTCACCATTTGAGACAA 4765

RESULT 5
LOCUS A22779 943 bp DNA linear PAT 24-JAN-1995
DEFINITION non-structural coding region.
ACCESSION A22779
VERSION A22779.1 GI:832940
KEYWORDS
SOURCE Hepatitis C virus
ORGANISM Hepatitis C virus
REFERENCE 1 (bases 1 to 943)
AUTHORS Brechot,C., Kreamsdorf,D. and Porchon,C.
TITLE Nucleotide and peptide sequences of an isolate of the hepatitis C
JOURNAL virus; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Hepacivirus.
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Db 630 GCGCGGCTTCTGGCTGTTGGCGCGGTATTGCGTATCCACAGGCTGCGTGCATAGTA 689
QY 1975 GG 1976

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Db 690 GG 691

RESULT 6
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DEFINITION Sequence 6 from patent US 5866139.
ACCESSION AR031209
VERSION AR031209.1 GI:5945498
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 943)
AUTHORS Brechot,C., Kreamsdorf,D. and Porchon,C.
TITLE Nucleotide and peptide sequences of a hepatitis C virus isolate,
JOURNAL diagnostic and therapeutic applications
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Db 630 GCGCGGCTTCTGGCTGTTGGCGCGGTATTGCGTATCCACAGGCTGCGTGCATAGTA 689
QY 1975 GG 1976
Db 690 GG 691

RESULT 7
LOCUS AR145025 943 bp DNA linear PAT 08-AUG-2001
DEFINITION Sequence 6 from patent US 6210962.
ACCESSION AR145025
VERSION AR145025.1 GI:15106892
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 943)
AUTHORS Brechot,C., Kreamsdorf,D. and Porchon,C.
TITLE Nucleotide and peptide sequences of an isolate of the hepatitis C
JOURNAL virus; diagnostic and therapeutic applications thereof
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Db 630 GCGCGGCTTCTGGCTGTTGGCGCGGTATTGCGTATCCACAGGCTGCGTGCATAGTA 689
QY 1975 GG 1976
Db 690 GG 691

RESULT 8

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HPCNS3NS4  
 LOCUS HPCNS3NS4 943 bp RNA linear VRL 02-FEB-1999  
 DEFINITION Hepatitis C virus NS3/NS4 gene for ORF 1, partial cds.  
 ACCESSION D10664 D01103  
 VERSION D10664.1 GI:221623  
 KEYWORDS nonstructural protein; NS3/NS4.  
 SOURCE Hepatitis C virus  
 ORGANISM Hepatitis C virus  
 Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae; Hepacivirus.  
 REFERENCE 1 (bases 1 to 943)  
 AUTHORS Kremsdorf, D., Porchon, C., Kim, J.P., Reyes, G.R. and Brechot, C.  
 TITLE Partial nucleotide sequence analysis of a French hepatitis C virus: implications for HCV genetic variability in the E2/NS1 protein  
 JOURNAL J. Gen. Virol. 72 (Pt 10), 2557-2561 (1991)  
 MEDLINE 92013977  
 PUBMED 1655961  
 REFERENCE 2 (sites)  
 AUTHORS Mink, M.A., Benichou, S., Madaule, P., Tiollais, P., Prince, A.M. and Inchausti, G.  
 TITLE Characterization and mapping of a B-cell immunogenic domain in hepatitis C virus E2 glycoprotein using a yeast peptide library  
 JOURNAL Virology 200 (1), 246-255 (1994)  
 MEDLINE 94174722  
 PUBMED 7510436  
 REFERENCE 3 (bases 1 to 943)  
 AUTHORS Kremsdorf, D.  
 TITLE Direct Submission  
 JOURNAL Submitted (17-JUL-1991) Dina Kremsdorf, INSERM U75; CHU Necker, 156, rue de Vaugirard, Paris 75015, France (E-mail: BEAUNESPRCITI51, Tel:40659911)  
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 LOCUS Hepatitis C virus isolate colonel complete genome.  
 DEFINITION AF290978  
 ACCESSION AF290978  
 VERSION AF290978.1 GI:9930556

KEYWORDS  
 SOURCE  
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 Hepatitis C virus  
 Hepatitis C virus  
 Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae; Hepacivirus.  
 REFERENCE 1 (bases 1 to 9365)  
 AUTHORS Desai, S.M., Devare, S. and Yamaguchi, J.  
 TITLE Hepatitis C virus  
 JOURNAL Unpublished  
 REFERENCE 2 (bases 1 to 9365)  
 AUTHORS Desai, S.M., Devare, S. and Yamaguchi, J.  
 TITLE Direct Submission  
 JOURNAL Submitted (28-JUL-2000) 90D, Abbott Laboratories, 1401 Sheridan Rd, North Chicago, IL 60064, USA  
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LTCITKAKACRAAGLQRTMLVCGDDLVVVCESAGVOEDAASLRFAETAMTRYSAAP  
GDQPEYDELELITSCSNVSVHAGDKRKYVYLTTRDPTPLARAWEATRHPTVNSW  
LGNIMEFAPTLWARMILMTHFFSVLIARDQPEQALNCEIYGCACVIEPLDPTIQRSL  
HGLSAFSLHSYSGEINRVAACLRKIGVPLRAWKHRARSVRARLLSRGGRALCGKY  
LFNNAVRTKPKLPIIAAGRLDLSGWTAGYSGGDIYHSVSHARPNFWCLLLILAAG  
VGIYLLPNR"

ORIGIN  
Query Match 3.0%; Score 62; DB 14; Length 9365;  
Best Local Similarity 100.0%; Pred. No. 2.1e-25;  
Matches 62; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 970 GCTACCCCTCCGGCTCGTCACTGTGCCCCCATCTTAACATCGAGAGGTTCCTCTCC 1029  
|||  
Db 4374 GCTACCCCTCCGGCTCGTCACTGTGCCCCCATCTTAACATCGAGAGGTTCCTCTCC 4433  
|||  
QY 1030 AC 1031  
|||  
Db 4434 AC 4435

RESULT 10  
AF511950 9395 bp RNA linear VRL 13-JUN-2002  
LOCUS Hepatitis C virus isolate XF224 polyprotein gene, complete  
DEFINITION  
ACCESSION AF511950  
VERSION AF511950.1 GI:21397077  
KEYWORDS  
SOURCE  
ORGANISM Hepatitis C virus  
Hepatitis C virus  
Hepatitis C virus  
Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;  
Hepacivirus

REFERENCE 1 (bases 1 to 9395)  
AUTHORS Fan, X. and Di Bisceglie, A.M.  
TITLE Clonal Nature of Hepatitis C Virus  
JOURNAL Unpublished  
REFERENCE 2 (bases 1 to 9395)  
AUTHORS Fan, X. and Di Bisceglie, A.M.  
TITLE Direct Submission  
JOURNAL Submitted (14-MAY-2002) Gastroenterology & Hepatology, Saint Louis University School of Medicine, 1402 South Grand Blvd., St. Louis, MO 63104, USA

FEATURES  
source  
1. .9395 Location/Qualifiers  
/organism="Hepatitis C virus"  
/mol\_type="genomic RNA"  
/strain="JLD6-2"  
/isolate="XF224"  
/db\_xref="taxon:11103"  
misc\_feature 1. .9395  
/note="similar to polyprotein"

ORIGIN  
Query Match 3.0%; Score 62; DB 14; Length 9395;  
Best Local Similarity 100.0%; Pred. No. 2.1e-25;  
Matches 62; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY -970 GCTACCCCTCCGGCTCGTCACTGTGCCCCCATCTTAACATCGAGAGGTTCCTCTCC 1029  
|||  
Db 4358 GCTACCCCTCCGGCTCGTCACTGTGCCCCCATCTTAACATCGAGAGGTTCCTCTCC 4417  
|||  
QY 1030 AC 1031  
|||  
Db 4418 AC 4419

RESULT 11  
BD226202 957 bp DNA linear PAT 17-JUL-2003  
LOCUS

DEFINITION Improved immunodiagnostic assays using reducing agents.  
ACCESSION BD226202  
VERSION BD226202.1 GI:33035972  
KEYWORDS JP 2002512370-A/4.  
SOURCE unidentifed  
ORGANISM unclassified  
unclassified.

REFERENCE 1 (bases 1 to 957)  
AUTHORS Maertens, G., Louwagie, J., Bosman, A., Sablon, E. and Zreïn, M.  
TITLE Improved immunodiagnostic assays using reducing agents  
JOURNAL Patent: JP 2002512370-A 4 23-APR-2002;  
INNOGENETICS NV

COMMENT OS Hepatitis virus (hepatitis C virus)  
PN JP 2002512370-A/4  
PD 23-APR-2002  
PF 15-APR-1999 JP 2000545027  
PR 17-APR-1998 EP 98870087.8  
PI GEERT MAERTENS, JOOST LOUWAGIE, ALFONS BOSMAN, ERWIN SABLON, MAAN ZREIN  
PC G01N33/543, C12N1/15, C12N1/19, C12N1/21, C12N5/10, C12N9/00 PC  
C12N15/09, C12Q1/25,  
PC C12Q1/68, G01N33/573, G01N33/576, C12N5/00, C12N15/00 CC  
Improved immunodiagnostic assays using reducing agents FH Key  
Location/Qualifiers  
FT source 1. .957  
virus), /organism="Hepatitis virus (hepatitis C FT

FEATURES  
source  
1. .957 Location/Qualifiers  
/organism="unidentified"  
/mol\_type="genomic DNA"  
/db\_xref="taxon:32644"

ORIGIN  
Query Match 3.0%; Score 61; DB 6; Length 957;  
Best Local Similarity 100.0%; Pred. No. 8.5e-25;  
Matches 61; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 710 TGGGCTTTCGTCTTACATGTCACAGGCCATCGGATTCATCTAACATCAGGACTGGG 769  
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Db 344 TGGGCTTTCGTCTTACATGTCACAGGCCATCGGATTCATCTAACATCAGGACTGGG 403  
|||  
QY 770 T 770  
|||  
Db 404 T 404

RESULT 12  
HPCNS10CLN 241 bp ss-RNA linear VRL 02-AUG-1993  
LOCUS Hepatitis C virus (clone #10) nonstructural protein (NS3/NS4) gene,  
DEFINITION partial cds.  
ACCESSION M94401 M84480  
VERSION M94401.1 GI:329772  
KEYWORDS nonstructural protein.  
SOURCE Hepatitis C virus  
ORGANISM Hepatitis C virus  
Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;  
Hepacivirus.

REFERENCE 1 (bases 1 to 241)  
AUTHORS Martell, M., Esteban, J.I., Quer, J., Genesca, J., Weiner, A., Esteban, R., Guardia, J. and Gomez, J.  
TITLE Hepatitis C virus (HCV) circulates as a population of different but closely related genomes: quasispecies nature of HCV genome distribution  
JOURNAL J. Virol. 66 (5), 3225-3229 (1992)  
MEDLINE 92219420  
PUBMED 1313927

COMMENT Original source text: Hepatitis C virus RNA.  
FEATURES  
1. .241 Location/Qualifiers  
/organism="Hepatitis C virus"  
/mol\_type="genomic RNA"

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/db xref="taxon:11103"
1. .241
/genes="NS3/NS4"
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/genes="NS3/NS4"
/codon_start=2
/product="nonstructural protein"
/protein_id="AAA45608.1"
/db xref="GI:329773"
/translation="LRAVMNTPGLPVCDHLEFWEVFTGLTHIDAHFLSQTQSGEN
LPLVAYQATVCARAQAQPPSWDQWVKLRLKFTL"

ORIGIN
Query Match 2.9%; Score 59; DB 14; Length 241;
Best Local Similarity 100.0%; Pred. No. 1.4e-23;
Matches 59; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1549 TACATGAACACCCCGGACTTCCCGTGTGCCAAGACCATCTTGATTTGGAGGGCGT 1607
DB 11 TACATGAACACCCCGGACTTCCCGTGTGCCAAGACCATCTTGATTTGGAGGGCGT 69

RESULT 13
HPCNS11CLN 241 bp ss-RNA linear VRL 02-AUG-1993
LOCUS Hepatitis C virus (clone #11) nonstructural protein (NS3/NS4) gene,
DEFINITION partial cds.
ACCESSION M94400.1 GI:329774
VERSION nonstructural protein.
KEYWORDS Hepatitis C virus
SOURCE Hepatitis C virus
ORGANISM Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Hepacivirus.
REFERENCE 1 (bases 1 to 241)
AUTHORS Martell,M., Esceban,J.I., Quer,J., Genesca,J., Weiner,A.,
Esteban,R., Guardia,J. and Gomez,J.
TITLE Hepatitis C virus (HCV) circulates as a population of different but
closely related genomes: quasisppecies nature of HCV genome
distribution
JOURNAL J. Virol. 66 (5), 3225-3229 (1992)
MEDLINE 92219420
PUBMED 1313927
COMMENT Original source text: Hepatitis C virus RNA.
FEATURES
source
1. .241
/organism="Hepatitis C virus"
/mol_type="genomic RNA"
/db_xref="taxon:11103"
1. .241
/genes="NS3/NS4"
<1. .>241
/genes="NS3/NS4"
/codon_start=2
/product="nonstructural protein"
/protein_id="AAA45609.1"
/db_xref="GI:329775"
/translation="LRAVMNTPGLPVCDHLEFWEVFTGLTHIDAHFLSQTQSGEN
LPLVAYQATVCARAQAQPPSWDQWVKLRLKFTL"

ORIGIN
Query Match 2.9%; Score 59; DB 14; Length 241;
Best Local Similarity 100.0%; Pred. No. 1.4e-23;
Matches 59; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1549 TACATGAACACCCCGGACTTCCCGTGTGCCAAGACCATCTTGATTTGGAGGGCGT 1607
DB 11 TACATGAACACCCCGGACTTCCCGTGTGCCAAGACCATCTTGATTTGGAGGGCGT 69

RESULT 14
HCVNSTP 550 bp RNA linear VRL 27-APR-1994
LOCUS Hepatitis C virus (strain Fla) plasma DNA.

```

```

DEFINITION Hepatitis C Virus sequence for non-structural protein.
ACCESSION X71406
VERSION 1 GI:296161
KEYWORDS hepatitis; non-structural protein; NS3/NS4.
SOURCE Hepatitis C virus
ORGANISM Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Hepacivirus.
REFERENCE 1
AUTHORS Vassilev,V.B., Viazov,S.O., Kotova,E.Y. and Mosikov,V.V.
TITLE Determination of the nucleotide sequence of the Russian variant of
the hepatitis C virus
JOURNAL Mol. Gen. Mikrobiol. Virusol. 1, 33-37 (1994)
AUTHORS Vassilev,V.B.
TITLE Direct Submission
JOURNAL Submitted (13-APR-1993) V.B. Vassilev, Institute of Genetics &
Selection of Industrial Microorganisms, 1-st Dorozhny proezd str.,
1, 113545 Moscow, Russia, USSR
FEATURES
source
1. .550
/organism="Hepatitis C virus"
/mol_type="genomic RNA"
/variety="Russian isolate"
/isolate="patient 11, with chronic non-A, non-B
post-transfusion hepatitis"
/db_xref="taxon:11103"
1. .550
/genes="NS3 /NS4"
<1. .>550
/genes="NS3 /NS4"
/codon_start=3
/product="non-structural protein"
/protein_id="C2M50530.1"
/db_xref="GI:296162"
1. .550
/db_xref="GOA:Q68960"
/db_xref="SPTREMBL:Q68960"
/translation="TVDFSLDPTFTTITTLPODASVTSQRRTGRGKPGIYRFVAP
GERPSMFDSVLCEYDAGCAWELTPTAETVRLRAYMNTPLPVCQHLSEWGVF
TGLTQIDAHLFSOTKSGENLPLVAYQATVCARAQAQPPSWDQWVKLRLKFTLHG
PTPLLYRLGAVQNEVTLTHPVT"

ORIGIN
Query Match 2.9%; Score 59; DB 14; Length 550;
Best Local Similarity 100.0%; Pred. No. 1.5e-23;
Matches 59; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1567 CTTCCTGGTGTGCCAAGACCATCTTGATTTGGAGGGCGTCTTTACGGTCTCACCCA 1625
DB 264 CTTCCTGGTGTGCCAAGACCATCTTGATTTGGAGGGCGTCTTTACGGTCTCACCCA 322

RESULT 15
HPCNS34 1477 bp DNA linear VRL 21-AUG-1995
LOCUS Hepatitis C virus nonstructural region.
DEFINITION Hepatitis C virus nonstructural region.
ACCESSION M60220
VERSION M60220.1 GI:329802
KEYWORDS nonstructural region.
SOURCE Hepatitis C virus
ORGANISM Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Hepacivirus.
REFERENCE 1 (bases 1 to 1477)
AUTHORS Li,J.S., Tong,S.P., Vitvitski,L., Lepot,D. and Trepo,C.
TITLE Two French genotypes of hepatitis C virus: homology of the
predominant genotype with the prototype American strain
JOURNAL Gene 105 (2), 167-172 (1991)
MEDLINE 92039028
PUBMED 1718820
COMMENT Original source text: Hepatitis C virus (strain Fla) plasma DNA.
FEATURES
source
1. .1477

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Db 33 ACCTGTGTCACCCAGACAGTCGACTTCAGCCTTGACCCCTTACCTTACCATTTGAGACAA 90

RESULT 18  
ARI24772  
LOCUS AR124772 1414 bp DNA linear PAT 16-MAY-2001  
DEFINITION Sequence 51 from patent US 6172189.  
ACCESSION AR124772  
VERSION AR124772.1 GI:14110133  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 1414)  
AUTHORS Devare,S.G., Desai,S.M., Casey,J.M., Dailey,S.H., Dawson,G.J.,  
Gutierrez,R.A., Lesniewski,R.R., Stewart,J.L. and Rupprecht,K.R.  
TITLE Hepatitis C assay utilizing recombinant antigens  
JOURNAL Patent: US 6172189-A 51 09-JAN-2001;  
FEATURES Location/Qualifiers  
source 1..1414  
/organism="unknown"  
/mol\_type="unassigned DNA"

ORIGIN  
Query Match 2.8%; Score 58; DB 6; Length 1414;  
Best Local Similarity 100.0%; Pred. No. 6.1e-23;  
Matches 58; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1291 ACCTGTGTCACCCAGACAGTCGACTTCAGCCTTGACCCCTTACCATTTGAGACAA 1348  
|||||  
Db 22 ACCTGTGTCACCCAGACAGTCGACTTCAGCCTTGACCCCTTACCATTTGAGACAA 79  
|||||

RESULT 19  
ARI24772  
LOCUS AR353559 1414 bp DNA linear PAT 17-AUG-2003  
DEFINITION Sequence 51 from patent US 6593083.  
ACCESSION AR353559  
VERSION AR353559.1 GI:33759549  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 1414)  
AUTHORS Devare,S.G., Desai,S.M., Casey,J.M., Dailey,S.H., Dawson,G.J.,  
Gutierrez,R.A., Lesniewski,R.R., Stewart,J.L. and Rupprecht,K.R.  
TITLE Hepatitis C assay utilizing recombinant antigens  
JOURNAL Patent: US 6593083-A 51 15-JUL-2003;  
FEATURES Location/Qualifiers  
source 1..1414  
/organism="unknown"  
/mol\_type="genomic DNA"

ORIGIN  
Query Match 2.8%; Score 58; DB 6; Length 1414;  
Best Local Similarity 100.0%; Pred. No. 6.1e-23;  
Matches 58; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1291 ACCTGTGTCACCCAGACAGTCGACTTCAGCCTTGACCCCTTACCATTTGAGACAA 1348  
|||||  
Db 22 ACCTGTGTCACCCAGACAGTCGACTTCAGCCTTGACCCCTTACCATTTGAGACAA 79  
|||||

RESULT 20  
ARI24774  
LOCUS AR124774 1420 bp DNA linear PAT 16-MAY-2001  
DEFINITION Sequence 57 from patent US 6172189.  
ACCESSION AR124774  
VERSION AR124774.1 GI:14110135  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.

/organism="Hepatitis C virus"  
/mol\_type="genomic DNA"  
/strain="Fla"  
/db\_xref="taxon:11103"  
/tissue\_type="plasma"

ORIGIN  
Query Match 2.9%; Score 59; DB 14; Length 1477;  
Best Local Similarity 100.0%; Pred. No. 1.5e-23;  
Matches 59; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1903 TGGGTGTCGTTGGCGGCTTCTGGCTGCTTGGCCGCGTATTCCTATCCACAGGCTG 1961  
632 TGGGTGTCGTTGGCGGCTTCTGGCTGCTTGGCCGCGTATTCCTATCCACAGGCTG 690  
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Db

RESULT 16  
ARI24773  
LOCUS AR124773 382 bp DNA linear PAT 16-MAY-2001  
DEFINITION Sequence 56 from patent US 6172189.  
ACCESSION AR124773  
VERSION AR124773.1 GI:14110134  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 382)  
AUTHORS Devare,S.G., Desai,S.M., Casey,J.M., Dailey,S.H., Dawson,G.J.,  
Gutierrez,R.A., Lesniewski,R.R., Stewart,J.L. and Rupprecht,K.R.  
TITLE Hepatitis C assay utilizing recombinant antigens  
JOURNAL Patent: US 6172189-A 56 09-JAN-2001;  
FEATURES Location/Qualifiers  
source 1..382  
/organism="unknown"  
/mol\_type="unassigned DNA"

ORIGIN  
Query Match 2.8%; Score 58; DB 6; Length 382;  
Best Local Similarity 100.0%; Pred. No. 6e-23;  
Matches 58; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1291 ACCTGTGTCACCCAGACAGTCGACTTCAGCCTTGACCCCTTACCATTTGAGACAA 1348  
|||||  
Db 33 ACCTGTGTCACCCAGACAGTCGACTTCAGCCTTGACCCCTTACCATTTGAGACAA 90  
|||||

RESULT 17  
ARI24774  
LOCUS AR353560 382 bp DNA linear PAT 17-AUG-2003  
DEFINITION Sequence 56 from patent US 6593083.  
ACCESSION AR353560  
VERSION AR353560.1 GI:33759550  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 382)  
AUTHORS Devare,S.G., Desai,S.M., Casey,J.M., Dailey,S.H., Dawson,G.J.,  
Gutierrez,R.A., Lesniewski,R.R., Stewart,J.L. and Rupprecht,K.R.  
TITLE Hepatitis C assay utilizing recombinant antigens  
JOURNAL Patent: US 6593083-A 56 15-JUL-2003;  
FEATURES Location/Qualifiers  
source 1..382  
/organism="unknown"  
/mol\_type="genomic DNA"

ORIGIN  
Query Match 2.8%; Score 58; DB 6; Length 382;  
Best Local Similarity 100.0%; Pred. No. 6e-23;  
Matches 58; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1291 ACCTGTGTCACCCAGACAGTCGACTTCAGCCTTGACCCCTTACCATTTGAGACAA 1348  
|||||

REFERENCE 1 (bases 1 to 1420)  
 Devare,S.G., Desai,S.M., Casey,J.M., Dailey,S.H., Dawson,G.J.,  
 Gutierrez,R.A., Lesniewski,R.R., Stewart,J.L. and Rupprecht,K.R.  
 TITLE Hepatitis C assay utilizing recombinant antigens  
 JOURNAL Patent: US 6172189-A 57 09-JAN-2001;  
 FEATURES Location/Qualifiers  
 source 1..1420  
 /organism="unknown"  
 /mol\_type="unassigned DNA"

ORIGIN

Query Match 2.8%; Score 58; DB 6; Length 1420;  
 Best Local Similarity 100.0%; Pred. No. 6.1e-23;  
 Matches 58; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1291 ACGTGTGTACCCAGACAGTCGACTTCAGCCTTGACCTTACCTTCACCATGAGCAA 1348  
 Db 22 ACGTGTGTACCCAGACAGTCGACTTCAGCCTTGACCTTACCTTCACCATGAGCAA 79

RESULT 21  
 LOCUS AR353561 1420 bp DNA linear PAT 17-AUG-2003  
 DEFINITION Sequence 57 from patent US 6593083  
 ACCESSION AR353561  
 VERSION AR353561.1 GI:33759551  
 KEYWORDS  
 SOURCE Unknown.  
 ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 1420)  
 Devare,S.G., Desai,S.M., Casey,J.M., Dailey,S.H., Dawson,G.J.,  
 Gutierrez,R.A., Lesniewski,R.R., Stewart,J.L. and Rupprecht,K.R.  
 TITLE Hepatitis C assay utilizing recombinant antigens  
 JOURNAL Patent: US 6593083-A 57 15-JUL-2003;  
 FEATURES Location/Qualifiers  
 source 1..1420  
 /organism="unknown"  
 /mol\_type="genomic DNA"

ORIGIN

Query Match 2.8%; Score 58; DB 6; Length 1420;  
 Best Local Similarity 100.0%; Pred. No. 6.1e-23;  
 Matches 58; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1291 ACGTGTGTACCCAGACAGTCGACTTCAGCCTTGACCTTACCTTCACCATGAGCAA 1348  
 Db 22 ACGTGTGTACCCAGACAGTCGACTTCAGCCTTGACCTTACCTTCACCATGAGCAA 79

RESULT 22  
 LOCUS HPCHCV35 162 bp ss-RNA linear VRL 20-MAY-1994  
 DEFINITION Hepatitis C virus mRNA, partial cds.  
 ACCESSION M55151  
 VERSION M55151.1 GI:329764  
 KEYWORDS  
 SOURCE Hepatitis C virus  
 ORGANISM Hepatitis C virus

REFERENCE 1 (bases 1 to 162)  
 Ulrich,P.P., Romeo,J.M., Lane,P.K., Kelly,I., Daniel,L.J. and  
 Vyas,G.N.  
 TITLE Detection, semiquantitation, and genetic variation in hepatitis C  
 virus sequences amplified from the plasma of blood donors with  
 elevated alanine aminotransferase  
 J. Clin. Invest. 86 (5), 1609-1614 (1990)  
 91056164  
 2173725  
 COMMENT Original source text: Hepatitis C virus, cDNA to genomic RNA, from  
 human plasma.  
 FEATURES Location/Qualifiers

1..162  
 /organism="Hepatitis C virus"  
 /mol\_type="genomic RNA"  
 /db\_xref="taxon:11103"  
 /map="2895-3056"  
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 /note="ORF"  
 /codon\_start=1  
 /protein\_id="AAA45605.1"  
 /db\_xref="GI:329765"  
 /translation="VAYYRGLDVSVIPTSGDVVVVATDALMTGYTGFDSVIDCNTCV  
 TQVDFSLDP"

ORIGIN

Query Match 2.8%; Score 57; DB 14; Length 162;  
 Best Local Similarity 100.0%; Pred. No. 2.5e-22;  
 Matches 57; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1273 TCGGTGATAGACTGCAACACGCTGTGTACCCAGACAGTCGACTTCAGCCTTGACCTT 1329  
 Db 106 TCGGTGATAGACTGCAACACGCTGTGTACCCAGACAGTCGACTTCAGCCTTGACCTT 162

RESULT 23

LOCUS HPCNS17CLN 223 bp ss-RNA linear VRL 02-AUG-1993  
 DEFINITION Hepatitis C virus (clone #17) nonstructural protein (NS3/NS4) gene,  
 partial cds.  
 ACCESSION M94451 M84480  
 VERSION M94451.1 GI:329778  
 KEYWORDS nonstructural protein.  
 SOURCE Hepatitis C virus  
 ORGANISM Hepatitis C virus

REFERENCE 1 (bases 1 to 223)  
 Martell,M., Esteban,J.I., Quer,J., Genesca,J., Weiner,A.,  
 Esteban,R., Guardia,J. and Gomez,J.  
 TITLE Hepatitis C virus (HCV) circulates as a population of different but  
 closely related genomes: Quasispecies nature of HCV genome  
 distribution  
 J. Virol. 66 (5), 3225-3229 (1992)  
 92219420  
 1313927  
 COMMENT Original source text: Hepatitis C virus RNA.  
 FEATURES Location/Qualifiers

ORIGIN

1..223  
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 /mol\_type="genomic RNA"  
 /db\_xref="taxon:11103"  
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 /gene="NS3/NS4"  
 /codon\_start=2  
 /product="nonstructural protein"  
 /protein\_id="AAA45611.1"  
 /db\_xref="GI:329779"  
 /translation="TPGLPVQCVHLEFWEGVFTGITHDAHFLSOTKSGENLPYLVA  
 YQATVCARAQAAPPSPWDQWKLRLKPTL"

ORIGIN

Query Match 2.8%; Score 57; DB 14; Length 223;  
 Best Local Similarity 100.0%; Pred. No. 2.5e-22;  
 Matches 57; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1723 CCTCCCCCGTCGTGGAGCAGATGCGAGTCTTGATCGTCTCAAGCCACCTTC 1779  
 Db 167 CCTCCCCCGTCGTGGAGCAGATGCGAGTCTTGATCGTCTTGATCGTCTCAAGCCACCTTC 228



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RESULT 24
HPCNS15CLN 229 bp ss-RNA linear VRL 02-AUG-1993
LOCUS Hepatitis C virus (clone #15) nonstructural protein (NS3/NS4) gene,
DEFINITION partial cds.
ACCESSION M94449 M84480
VERSION M94449.1 GI:329776
KEYWORDS nonstructural protein.
SOURCE Hepatitis C virus
ORGANISM Hepatitis C virus
Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Hepacivirus.
REFERENCE 1 (bases 1 to 229)
AUTHORS Martell,M., Esteban,J.I., Quer,J., Genesca,J., Weiner,A.,
Esteban,R., Guardia,J. and Gomez,J.
TITLE Hepatitis C virus (HCV) circulates as a population of different but
closely related genomes: quasisppecies nature of HCV genome
distribution
J. Virol. 66 (5), 3225-3229 (1992)
JOURNAL 92219420
MEDLINE 1313927
PUBMED
COMMENT Original source text: Hepatitis C virus RNA.
FEATURES
source
1..229
Location/Qualifiers
1..229
/organism="Hepatitis C virus"
/mol_type="genomic RNA"
/db_xref="taxon:11103"
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/gene="NS3/NS4"
<1..>229
/gene="NS3/NS4"
/codon_start=2
/product="nonstructural protein"
/protein_id="AAA45610.1"
/db_xref="GI:329777"
/translation="MNTPLPVCODHLEFWGVFTGLTHIDAFSLQTKQSGENLPL"
VAYQATVCARAQAAPPSPWDQWKILRLKPTL"
ORIGIN
1551 CATGAACACCCGGGACTTCCCGTGTGCCAAGACCATCTTGAATTTTGGAGGCGT 1607
|||||
Db 1 CATGAACACCCGGGACTTCCCGTGTGCCAAGACCATCTTGAATTTTGGAGGCGT 57
|||||

Query Match 2.8%; Score 57; DB 14; Length 229;
Best Local Similarity 100.0%; Pred. No. 2.5e-22;
Matches 57; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1551 CATGAACACCCGGGACTTCCCGTGTGCCAAGACCATCTTGAATTTTGGAGGCGT 1607
|||||
Db 1 CATGAACACCCGGGACTTCCCGTGTGCCAAGACCATCTTGAATTTTGGAGGCGT 57
|||||

RESULT 25
HPCNSCLN5 241 bp ss-RNA linear VRL 02-AUG-1993
LOCUS Hepatitis C virus (clone #5) nonstructural protein (NS3/NS4) gene,
DEFINITION partial cds
ACCESSION M94469 M84480
VERSION M94469.1 GI:329852
KEYWORDS nonstructural protein.
SOURCE Hepatitis C virus
ORGANISM Hepatitis C virus
Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Hepacivirus.
REFERENCE 1 (bases 1 to 241)
AUTHORS Martell,M., Esteban,J.I., Quer,J., Genesca,J., Weiner,A.,
Esteban,R., Guardia,J. and Gomez,J.
TITLE Hepatitis C virus (HCV) circulates as a population of different but
closely related genomes: quasisppecies nature of HCV genome
distribution
J. Virol. 66 (5), 3225-3229 (1992)
JOURNAL 92219420
MEDLINE 1313927
PUBMED
COMMENT Original source text: Hepatitis C virus RNA.
FEATURES
source
1..241
Location/Qualifiers
1..241

Query Match 2.8%; Score 57; DB 14; Length 229;
Best Local Similarity 100.0%; Pred. No. 2.5e-22;
Matches 57; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1551 CATGAACACCCGGGACTTCCCGTGTGCCAAGACCATCTTGAATTTTGGAGGCGT 1607
|||||
Db 1 CATGAACACCCGGGACTTCCCGTGTGCCAAGACCATCTTGAATTTTGGAGGCGT 57
|||||

RESULT 26
HPCNS15CLN 281 bp DNA linear PAT 06-FEB-1997
LOCUS Hepatitis C virus (clone #15) nonstructural protein (NS3/NS4) gene,
DEFINITION partial cds
ACCESSION M94449 M84480
VERSION M94449.1 GI:329776
KEYWORDS nonstructural protein.
SOURCE Hepatitis C virus
ORGANISM Hepatitis C virus
Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Hepacivirus.
REFERENCE 1 (bases 1 to 281)
AUTHORS Martell,M., Esteban,J.I., Quer,J., Genesca,J., Weiner,A.,
Esteban,R., Guardia,J. and Gomez,J.
TITLE Hepatitis C virus (HCV) circulates as a population of different but
closely related genomes: quasisppecies nature of HCV genome
distribution
J. Virol. 66 (5), 3225-3229 (1992)
JOURNAL 92219420
MEDLINE 1313927
PUBMED
COMMENT Original source text: Hepatitis C virus RNA.
FEATURES
source
1..281
Location/Qualifiers
1..281
/organism="Hepatitis C virus"
/mol_type="genomic RNA"
/db_xref="taxon:11103"
1..281
/gene="NS3/NS4"
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/gene="NS3/NS4"
/codon_start=2
/product="nonstructural protein"
/protein_id="AAA45675.1"
/db_xref="GI:329853"
/translation="LPLVAYQATVCARAQAAPPSPWDQWKILRLKPTL"
LPLVAYQATVCARAQAAPPSPWDQWKILRLKPTL"
ORIGIN
1723 CTTCCCTCCCTGCTGGGACGACAGATGCGAGTCTTGAATCCGTCCTCAGCCACCTC 1779
|||||
Db 185 CTTCCCTCCCTGCTGGGACGACAGATGCGAGTCTTGAATCCGTCCTCAGCCACCTC 241
|||||

Query Match 2.8%; Score 57; DB 14; Length 241;
Best Local Similarity 100.0%; Pred. No. 2.5e-22;
Matches 57; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1723 CTTCCCTCCCTGCTGGGACGACAGATGCGAGTCTTGAATCCGTCCTCAGCCACCTC 1779
|||||
Db 185 CTTCCCTCCCTGCTGGGACGACAGATGCGAGTCTTGAATCCGTCCTCAGCCACCTC 241
|||||

RESULT 27
HPCNSCLN5 281 bp DNA linear PAT 06-FEB-1997
LOCUS Hepatitis C virus (clone #5) nonstructural protein (NS3/NS4) gene,
DEFINITION partial cds
ACCESSION M94469 M84480
VERSION M94469.1 GI:329852
KEYWORDS nonstructural protein.
SOURCE Hepatitis C virus
ORGANISM Hepatitis C virus
Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Hepacivirus.
REFERENCE 1 (bases 1 to 281)
AUTHORS Martell,M., Esteban,J.I., Quer,J., Genesca,J., Weiner,A.,
Esteban,R., Guardia,J. and Gomez,J.
TITLE Hepatitis C virus (HCV) circulates as a population of different but
closely related genomes: quasisppecies nature of HCV genome
distribution
J. Virol. 66 (5), 3225-3229 (1992)
JOURNAL 92219420
MEDLINE 1313927
PUBMED
COMMENT Original source text: Hepatitis C virus RNA.
FEATURES
source
1..281
Location/Qualifiers
1..281
/organism="Hepatitis C virus"
/mol_type="genomic RNA"
/db_xref="taxon:11103"
1..281
/gene="NS3/NS4"
<1..>281
/gene="NS3/NS4"
/codon_start=2
/product="nonstructural protein"
/protein_id="AAA45675.1"
/db_xref="GI:329853"
/translation="LPLVAYQATVCARAQAAPPSPWDQWKILRLKPTL"
LPLVAYQATVCARAQAAPPSPWDQWKILRLKPTL"
ORIGIN
1723 CTTCCCTCCCTGCTGGGACGACAGATGCGAGTCTTGAATCCGTCCTCAGCCACCTC 1779
|||||
Db 185 CTTCCCTCCCTGCTGGGACGACAGATGCGAGTCTTGAATCCGTCCTCAGCCACCTC 241
|||||

Query Match 2.7%; Score 56; DB 6; Length 281;
Best Local Similarity 100.0%; Pred. No. 1e-21;
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTGGGCTCTCCGACCTTTACCTGTCACGAGCAGCGGATGTCATTC 347
|||||
Db 4 TGCACCTTGGGCTCTCCGACCTTTACCTGTCACGAGCAGCGGATGTCATTC 59
|||||

RESULT 28
HPCNSCLN5 281 bp DNA linear PAT 06-FEB-1997
LOCUS Hepatitis C virus (clone #5) nonstructural protein (NS3/NS4) gene,
DEFINITION partial cds
ACCESSION M94469 M84480
VERSION M94469.1 GI:329852
KEYWORDS nonstructural protein.
SOURCE Hepatitis C virus
ORGANISM Hepatitis C virus
Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Hepacivirus.
REFERENCE 1 (bases 1 to 281)
AUTHORS Martell,M., Esteban,J.I., Quer,J., Genesca,J., Weiner,A.,
Esteban,R., Guardia,J. and Gomez,J.
TITLE Hepatitis C virus (HCV) circulates as a population of different but
closely related genomes: quasisppecies nature of HCV genome
distribution
J. Virol. 66 (5), 3225-3229 (1992)
JOURNAL 92219420
MEDLINE 1313927
PUBMED
COMMENT Original source text: Hepatitis C virus RNA.
FEATURES
source
1..281
Location/Qualifiers
1..281
/organism="Hepatitis C virus"
/mol_type="genomic RNA"
/db_xref="taxon:11103"
1..281
/gene="NS3/NS4"
<1..>281
/gene="NS3/NS4"
/codon_start=2
/product="nonstructural protein"
/protein_id="AAA45675.1"
/db_xref="GI:329853"
/translation="LPLVAYQATVCARAQAAPPSPWDQWKILRLKPTL"
LPLVAYQATVCARAQAAPPSPWDQWKILRLKPTL"
ORIGIN
1723 CTTCCCTCCCTGCTGGGACGACAGATGCGAGTCTTGAATCCGTCCTCAGCCACCTC 1779
|||||
Db 185 CTTCCCTCCCTGCTGGGACGACAGATGCGAGTCTTGAATCCGTCCTCAGCCACCTC 241
|||||
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Query Match 2.7%; Score 56; DB 6; Length 281;  
 Best Local Similarity 100.0%; Pred. No. 1e-21;  
 Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTGGCGCTCCTCGGACCTTTACTGTCACGAGGCACGCCGATGTCATTC 347  
 Db 4 TGCACCTTGGCGCTCCTCGGACCTTTACTGTCACGAGGCACGCCGATGTCATTC 59

RESULT 28  
 LOCUS I82486 281 bp DNA linear PAT 10-JUN-1998  
 DEFINITION Sequence 75 from patent US 5712145.  
 ACCESSION I82486  
 VERSION I82486.1 GI:3210783  
 KEYWORDS  
 SOURCE Unknown.  
 ORGANISM Unknown.  
 REFERENCE 1 (bases 1 to 281)  
 AUTHORS Houghton, M., Choo, Q.-L. and Kuo, G.  
 TITLE Hepatitis C virus protease  
 JOURNAL Patent: US 5712145-A 75 27-JAN-1998;  
 FEATURES  
 source 1..281  
 /organism="unknown"  
 /mol\_type="unassigned DNA"

ORIGIN

Query Match 2.7%; Score 56; DB 6; Length 281;  
 Best Local Similarity 100.0%; Pred. No. 1e-21;  
 Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTGGCGCTCCTCGGACCTTTACTGTCACGAGGCACGCCGATGTCATTC 347  
 Db 4 TGCACCTTGGCGCTCCTCGGACCTTTACTGTCACGAGGCACGCCGATGTCATTC 59

RESULT 29  
 LOCUS BD140411 281 bp DNA linear PAT 18-SEP-2002  
 DEFINITION Hepatitis C virus protease.  
 ACCESSION BD140411  
 VERSION BD140411.1 GI:23235356  
 KEYWORDS JP 2002051791-A/29.  
 SOURCE Hepatitis C virus  
 ORGANISM Hepatitis C virus  
 REFERENCE 1 (bases 1 to 281)  
 AUTHORS Houghton, M., Choo, Q.-L. and Kuo, G.  
 TITLE Hepatitis C virus protease  
 JOURNAL Patent: JP 2002051791-A 29 19-FEB-2002;  
 COMMENT CHIRON CORP  
 OS HCV  
 PN JP 2002051791-A/29  
 PD 19-FEB-2002  
 PR 11-JUN-2001 JP 2001176369  
 PF 04-APR-1990 US 505433  
 PI MICHAEL HOUGHTON, QUI LIM CHOO, GEORGE KUO  
 PC C12N15/09, C12N15/09, C12N9/50, C12R1/93, (C12N15/09,  
 PC C12R1/93), C12N15/00, (C12N15/00, C12R1/93)  
 PC C12N15/00, C12N15/00, (C12N15/00, C12R1/93)  
 CC Hepatitis C virus protease  
 FH Key Location/Qualifiers  
 FT CDS  
 FEATURES  
 source 1..281  
 /organism="Hepatitis C virus"  
 /mol\_type="genomic DNA"  
 /db\_xref="taxon:11103"

ORIGIN

Query Match 2.7%; Score 56; DB 6; Length 281;  
 Best Local Similarity 100.0%; Pred. No. 1e-21;  
 Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTGGCGCTCCTCGGACCTTTACTGTCACGAGGCACGCCGATGTCATTC 347  
 Db 4 TGCACCTTGGCGCTCCTCGGACCTTTACTGTCACGAGGCACGCCGATGTCATTC 59

RESULT 30  
 LOCUS AR118676 283 bp DNA linear PAT 16-MAY-2001  
 DEFINITION Sequence 33 from patent US 6150087.  
 ACCESSION AR118676  
 VERSION AR118676.1 GI:14100586  
 KEYWORDS  
 SOURCE Unknown.  
 ORGANISM Unknown.  
 REFERENCE 1 (bases 1 to 283)  
 AUTHORS Chien, D.Y.  
 TITLE NANV diagnostics and vaccines  
 JOURNAL Patent: US 6150087-A 33 21-NOV-2000;  
 FEATURES  
 Location/Qualifiers  
 source 1..283  
 /organism="unknown"  
 /mol\_type="unassigned DNA"

ORIGIN

Query Match 2.7%; Score 56; DB 6; Length 283;  
 Best Local Similarity 100.0%; Pred. No. 1e-21;  
 Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTGGCGCTCCTCGGACCTTTACTGTCACGAGGCACGCCGATGTCATTC 347  
 Db 6 TGCACCTTGGCGCTCCTCGGACCTTTACTGTCACGAGGCACGCCGATGTCATTC 61

RESULT 31  
 LOCUS I32188 368 bp DNA linear PAT 06-FEB-1997  
 DEFINITION Sequence 71 from patent US 5585258.  
 ACCESSION I32188  
 VERSION I32188.1 GI:1822979  
 KEYWORDS  
 SOURCE Unknown.  
 ORGANISM Unknown.  
 REFERENCE 1 (bases 1 to 368)  
 AUTHORS Houghton, M., Choo, Q.-L. and Kuo, G.  
 TITLE Hepatitis C virus protease  
 JOURNAL Patent: US 5585258-A 71 17-DEC-1996;  
 FEATURES  
 Location/Qualifiers  
 source 1..368  
 /organism="unknown"  
 /mol\_type="unassigned DNA"

ORIGIN

Query Match 2.7%; Score 56; DB 6; Length 368;  
 Best Local Similarity 100.0%; Pred. No. 1e-21;  
 Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTGGCGCTCCTCGGACCTTTACTGTCACGAGGCACGCCGATGTCATTC 347  
 Db 220 TGCACCTTGGCGCTCCTCGGACCTTTACTGTCACGAGGCACGCCGATGTCATTC 275

RESULT 32  
 LOCUS I34279 368 bp DNA linear PAT 06-FEB-1997  
 DEFINITION Sequence 71 from patent US 5597691.  
 ACCESSION I34279  
 VERSION I34279.1 GI:1825070

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KEYWORDS
SOURCE      Unknown.
ORGANISM     Unknown.
REFERENCE    Unclassified.
1 (bases 1 to 368)
AUTHORS      Houghton,M., Choo,Q.-L. and Kuo,G.
TITLE        Hepatitis C virus protease
JOURNAL      Patent: US 5597691-A 71 28-JAN-1997;
FEATURES
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    1..368
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    /mol_type="unassigned DNA"
ORIGIN
Query Match      2.7%; Score 56; DB 6; Length 368;
Best Local Similarity 100.0%; Pred. No. 1e-21;
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 292 TGCACCTTGGCGCTCCTCGGACCTTTACCTGGTACAGGACGCGCGATGTCATTC 347
Db 220 TGCACCTTGGCGCTCCTCGGACCTTTACCTGGTACAGGACGCGCGATGTCATTC 275

RESULT 33
182484
LOCUS       I82484               368 bp      DNA      linear      PAT 10-JUN-1998
DEFINITION Sequence 71 from patent US 5712145.
ACCESSION   I82484
VERSION     I82484.1 GI:3210781
KEYWORDS    Unknown.
SOURCE      Unknown.
ORGANISM     Unclassified.
1 (bases 1 to 368)
AUTHORS      Houghton,M., Choo,Q.-L. and Kuo,G.
TITLE        Hepatitis C virus protease
JOURNAL      Patent: US 5712145-A 71 27-JAN-1998;
FEATURES
  source
    1..368
    /organism="unknown"
    /mol_type="unassigned DNA"
ORIGIN
Query Match      2.7%; Score 56; DB 6; Length 368;
Best Local Similarity 100.0%; Pred. No. 1e-21;
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 292 TGCACCTTGGCGCTCCTCGGACCTTTACCTGGTACAGGACGCGCGATGTCATTC 347
Db 220 TGCACCTTGGCGCTCCTCGGACCTTTACCTGGTACAGGACGCGCGATGTCATTC 275

RESULT 34
BD140409
LOCUS       BD140409             368 bp      DNA      linear      PAT 18-SEP-2002
DEFINITION Hepatitis C virus protease.
ACCESSION   BD140409
VERSION     BD140409.1 GI:23235354
KEYWORDS    JP 2002051791-A/27.
SOURCE      Hepatitis C virus
ORGANISM     Hepatitis C virus
1 (bases 1 to 368)
AUTHORS      Houghton,M., Choo,Q.-L. and Kuo,G.
TITLE        Hepatitis C virus protease
JOURNAL      Patent: JP 2002051791-A 27 19-FEB-2002;-
COMMENT      CHIRON CORP
OS          HCV
PN          JP 2002051791-A/27
PD          19-FEB-2002
PF          11-JUN-2001 JP 2001176369
PR          04-APR-1990 US 505433

KEYWORDS
SOURCE      Unknown.
ORGANISM     Unknown.
REFERENCE    Unclassified.
1 (bases 1 to 368)
AUTHORS      Houghton,M., Choo,Q.-L. and Kuo,G.
TITLE        Hepatitis C virus protease
JOURNAL      Patent: US 5597691-A 71 28-JAN-1997;
FEATURES
  source
    1..368
    /organism="unknown"
    /mol_type="unassigned DNA"
ORIGIN
Query Match      2.7%; Score 56; DB 6; Length 368;
Best Local Similarity 100.0%; Pred. No. 1e-21;
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 292 TGCACCTTGGCGCTCCTCGGACCTTTACCTGGTACAGGACGCGCGATGTCATTC 347
Db 220 TGCACCTTGGCGCTCCTCGGACCTTTACCTGGTACAGGACGCGCGATGTCATTC 275

RESULT 35
AF369231
LOCUS       AF369231             543 bp      RNA      linear      VRL 03-JUL-2002
DEFINITION Hepatitis C virus Pt.25 NS3 protease gene, partial cds.
ACCESSION   AF369231
VERSION     AF369231.1 GI:14150586
KEYWORDS    Hepatitis C virus
SOURCE      Hepatitis C virus
ORGANISM     Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
1 (bases 1 to 543)
AUTHORS      Holland-Staley,C.A., Kovari,L.C., Golenberg,E.M., Pobursky,K.J. and
Mayers,D.L.
TITLE        Genetic diversity and response to IFN of the NS3 protease gene from
clinical strains of the hepatitis C virus
JOURNAL      Arch. Virol. 147 (7), 1385-1406 (2002)
MEDLINE     22105140
PUBMED      12111414
REFERENCE    2 (bases 1 to 543)
AUTHORS      Holland-Staley,C.A., Kovari,L.C., Golenberg,E. and Mayers,D.L.
TITLE        Direct Submission
JOURNAL      Submitted (09-APR-2001) Infectious Disease Research, Henry Ford
Health Systems, 2799 W. Grand Blvd. Rm 7045 E & R, Detroit, MI
48202, USA
FEATURES
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    1..543
    /organism="Hepatitis C virus"
    /mol_type="genomic RNA"
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    /db_xref="taxon:11103"
    /note="type: 1A"
    <1..>543
    /note="polyprotein"
    /codon_start=1
    /product="NS3 protease"
    /protein_id="AAKS4556.1"
    /db_xref="GI:14150587"
    /translation="APITAYAAQTRGLGCIITSTIGRDKNQVEGOIVSTAAQTFL
ATCINGCVTVYHGAGTETLASPKGPVTQMTYVDQDLVGVAVQGAESLTECTGSS
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RGVAKAVDFIPVESLETTMS"
ORIGIN
Query Match      2.7%; Score 56; DB 14; Length 543;
Best Local Similarity 100.0%; Pred. No. 1e-21;
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 292 TGCACCTTGGCGCTCCTCGGACCTTTACCTGGTACAGGACGCGCGATGTCATTC 347
Db 220 TGCACCTTGGCGCTCCTCGGACCTTTACCTGGTACAGGACGCGCGATGTCATTC 275
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/db\_xref="RENTREMBL:CAD32154"  
 /translation="MAPITAYAOCTRGGLGCLILSLTGRDKNOVEGEVOIVSTAQTF  
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 SDLYLTVRHADVIPIVRERGSGRLSPFISYLVKSGGGLPCPGHAGVGIPIRAVC  
 TRGAKAVDFIPVENLETTMRSPVFTDINSPPVPPQSFVAHLHPTGSGSKTKVPA  
 YAAQGYKVLNLSVAATLFGAYMSKAHGINRTGRTVITITGSPITYSTYKFLA  
 DGGSGGAYDIIICDECHSDATSLIGITVLDQAGTAGARLVVLAATATPPGVTVP  
 PNIIEVALSTGPIPFYKAIPLKVIKGRHLIFCHSKKCDLAAKLVALGINAVAY  
 YRGLDVSIPPIGDVVVATDALMTGVTGDPDSVIDCNTCTVOTVDPGLDPTFTETI  
 TLQDASVTRORGRGKGLYRFVARGERSGMDSSVLCEYDAGCAYELTPA  
 ETVRLRAYNTPGLPVCQDHLFEWGVFTGLTHIDAHFLSQKSGENLPIYAVQA  
 TVCARAQAPPSWDMKCLIRKLPGLHGTPLLYRLGAVONEITLTHPVTKYITMCM  
 SADLEVVTSTWLVGGVLAALAYCLSTGCVIIVGRVLSGRPAIIPDREVLYREFDE  
 MEEC"

## ORIGIN

Query Match 2.7%; Score 56; DB 6; Length 2058;  
 Best Local Similarity 100.0%; Pred. No. 1.1e-21;  
 Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 292 TGCACTTGGCGCTCTCGGACCTTTACCTGTCACGAGGCACGCCGATGTCATTCC 347  
 Db 292 TGCACTTGGCGCTCTCGGACCTTTACCTGTCACGAGGCACGCCGATGTCATTCC 347

## RESULT 40

AX454818  
 LOCUS 2058 bp DNA linear PAT 06-JUL-2002  
 DEFINITION Sequence 1 from Patent WO196870.  
 ACCESSION AX454818

VERSION AX454818.1 GI:21714047

KEYWORDS synthetic construct

SOURCE synthetic construct

ORGANISM artificial sequences.

## REFERENCE

1 Chien,D.Y., Arcangel,P., Tandeske,L., George-Nascimento,C.,

Cicit,D. and Medina-Seiby,A.

Immunassays for anti-hcv antibodies

Patent: WO 0196870-A 1 20-DEC-2001;

CHIRON CORPORATION (US)

## FEATURES

source 1..2058

/organism="synthetic construct"

/mol\_type="unassigned DNA"

/db\_xref="taxon:32630"

/note="representative NS3/4a conformational antigen"

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/note="unnamed protein product"

/codon\_start=1

/transl\_table=11

/protein\_id="CAD38232.1"

/db\_xref="GI:21714048"

/db\_xref="RENTREMBL:CAD38232"

/translations="MAPITAYAOCTRGGLGCLILSLTGRDKNOVEGEVOIVSTAQTF  
 LATCINGCVTVHAGCTRIASPKGVIOYNTVNDQDLVGNWPAQGSRLTPTCTGS  
 SDLYLTVRHADVIPIVRERGSGRLSPFISYLVKSGGGLPCPGHAGVGIPIRAVC  
 TRGAKAVDFIPVENLETTMRSPVFTDINSPPVPPQSFVAHLHPTGSGSKTKVPA  
 YAAQGYKVLNLSVAATLFGAYMSKAHGINRTGRTVITITGSPITYSTYKFLA  
 DGGSGGAYDIIICDECHSDATSLIGITVLDQAGTAGARLVVLAATATPPGVTVP  
 PNIIEVALSTGPIPFYKAIPLKVIKGRHLIFCHSKKCDLAAKLVALGINAVAY  
 YRGLDVSIPPIGDVVVATDALMTGVTGDPDSVIDCNTCTVOTVDPGLDPTFTETI  
 TLQDASVTRORGRGKGLYRFVARGERSGMDSSVLCEYDAGCAYELTPA  
 ETVRLRAYNTPGLPVCQDHLFEWGVFTGLTHIDAHFLSQKSGENLPIYAVQA  
 TVCARAQAPPSWDMKCLIRKLPGLHGTPLLYRLGAVONEITLTHPVTKYITMCM  
 SADLEVVTSTWLVGGVLAALAYCLSTGCVIIVGRVLSGRPAIIPDREVLYREFDE  
 MEEC"

## ORIGIN

Query Match 2.7%; Score 56; DB 6; Length 2058;  
 Best Local Similarity 100.0%; Pred. No. 1.1e-21;  
 Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 292 TGCACTTGGCGCTCTCGGACCTTTACCTGTCACGAGGCACGCCGATGTCATTCC 347  
 Db 292 TGCACTTGGCGCTCTCGGACCTTTACCTGTCACGAGGCACGCCGATGTCATTCC 347

## RESULT 41

LOCUS 2064 bp DNA linear PAT 06-FEB-1997  
 DEFINITION Sequence 69 from patent US 5585258.  
 ACCESSION I32187

VERSION I32187.1 GI:1822978

KEYWORDS Unknown.

SOURCE Unknown.

ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 2064)

AUTHORS Houghton,M., Choo,Q.-L. and Kuo,G.

TITLE Hepatitis C virus protease

JOURNAL Patent: US 5585258-A 69 17-DEC-1996;

FEATURES Location/Qualifiers

source 1..2064

/organism="unknown"

/mol\_type="unassigned DNA"

## ORIGIN

Query Match 2.7%; Score 56; DB 6; Length 2064;

Best Local Similarity 100.0%; Pred. No. 1.1e-21;

Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 292 TGCACTTGGCGCTCTCGGACCTTTACCTGTCACGAGGCACGCCGATGTCATTCC 347  
 Db 541 TGCACTTGGCGCTCTCGGACCTTTACCTGTCACGAGGCACGCCGATGTCATTCC 596

## RESULT 42

LOCUS 2064 bp DNA linear PAT 06-FEB-1997  
 DEFINITION Sequence 69 from patent US 5597691-  
 ACCESSION I34278

VERSION I34278.1 GI:1825069

KEYWORDS Unknown.

SOURCE Unknown.

ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 2064)

AUTHORS Houghton,M., Choo,Q.-L. and Kuo,G.

TITLE Hepatitis C virus protease

JOURNAL Patent: US 5597691-A 69 28-JAN-1997;

FEATURES Location/Qualifiers

source 1..2064

/organism="unknown"

/mol\_type="unassigned DNA"

## ORIGIN

Query Match 2.7%; Score 56; DB 6; Length 2064;

Best Local Similarity 100.0%; Pred. No. 1.1e-21;

Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 292 TGCACTTGGCGCTCTCGGACCTTTACCTGTCACGAGGCACGCCGATGTCATTCC 347  
 Db 541 TGCACTTGGCGCTCTCGGACCTTTACCTGTCACGAGGCACGCCGATGTCATTCC 596

## RESULT 43

LOCUS 2064 bp DNA linear PAT 10-JUN-1998  
 DEFINITION Sequence 69 from patent US 5712145.  
 ACCESSION I82483

VERSION I82483.1 GI:3210780

KEYWORDS Unknown.

SOURCE Unknown.

ORGANISM Unclassified.

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REFERENCE 1 (bases 1 to 2064)
AUTHORS Houghton,M., Choo,Q.-L. and Kuo,G.
TITLE Hepatitis C virus protease
JOURNAL Patent: US 5712145-A 69 27-JAN-1998;
FEATURES Location/Qualifiers
source 1..2064
/organism="unknown"
/mol_type="unassigned DNA"

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Best Local Similarity 100.0%; Pred. No. 1.1e-21;
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTCGGCTCCTCGGACCTTACCTGTGTACAGGACGCGCGATGTCATTC 347
Db 541 TGCACCTTCGGCTCCTCGGACCTTACCTGTGTACAGGACGCGCGATGTCATTC 596

RESULT 44
BD140408 2064 bp DNA linear PAT 18-SEP-2002
LOCUS Hepatitis C virus protease.
DEFINITION BDI40408
ACCESSION BDI40408
VERSION BDI40408.1 GI:23235353
KEYWORDS Hepatitis C virus
SOURCE Hepatitis C virus
ORGANISM Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Hepacivirus.
REFERENCE 1 (bases 1 to 2064)
AUTHORS Houghton,M., Choo,Q.-L. and Kuo,G.
TITLE Hepatitis C virus protease
JOURNAL Patent: JP 2002051791-A 26 19-FEB-2002;
CHIRON CORP
COMMENT OS HCV
PN JP 2002051791-A/26
PD 19-FEB-2002
PF 11-JUN-2001 JP 2001176369
PR 04-APR-1990 US 505433
PI MICHAEL HOUGHTON,QUI LIM CHOO,GEORGE KUO
PC C12N15/09,C12N15/09,C12N9/50,C12R1/93),(C12N15/09,
PC C12R1/93),
PC C12N15/00,C12N15/00,(C12N15/00,C12R1/93)
CC Hepatitis C virus protease
FH Key Location/Qualifiers
FT CDS (7)..(2064).

FEATURES source
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Best Local Similarity 100.0%; Pred. No. 1.1e-21;
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTCGGCTCCTCGGACCTTACCTGTGTACAGGACGCGCGATGTCATTC 347
Db 541 TGCACCTTCGGCTCCTCGGACCTTACCTGTGTACAGGACGCGCGATGTCATTC 596

RESULT 45
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LOCUS Hepatitis C virus protease.
DEFINITION BDI40416
ACCESSION BDI40416
VERSION BDI40416.1 GI:23235361
KEYWORDS JP 2002051791-A/34..
SOURCE Hepatitis C virus
ORGANISM Viruses; ssRNA positive-strand viruses, no DNA stage; Flaviviridae;
Hepacivirus.

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Hepacivirus.
REFERENCE 1 (bases 1 to 2283)
AUTHORS Houghton,M., Choo,Q.-L. and Kuo,G.
TITLE Hepatitis C virus protease
JOURNAL Patent: JP 2002051791-A 34 19-FEB-2002;
CHIRON CORP
COMMENT OS HCV
PN JP 2002051791-A/34
PD 19-FEB-2002
PF 11-JUN-2001 JP 2001176369
PR 04-APR-1990 US 505433
PI MICHAEL HOUGHTON,QUI LIM CHOO,GEORGE KUO
PC C12N15/09,C12N15/09,C12N9/50,C12R1/93),(C12N15/09,
PC C12R1/93),
PC C12N15/00,C12N15/00,(C12N15/00,C12R1/93)
CC Hepatitis C virus protease
FH Key Location/Qualifiers
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Db 1000 TGCACCTTCGGCTCCTCGGACCTTACCTGTGTACAGGACGCGCGATGTCATTC 1055

RESULT 46
I32195 2523 bp DNA linear PAT 06-FEB-1997
LOCUS Sequence 85 from patent US 5585259.
DEFINITION I32195
ACCESSION I32195
VERSION I32195.1 GI:1822986
KEYWORDS SOURCE
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 2523)
AUTHORS Houghton,M., Choo,Q.-L. and Kuo,G.
TITLE Hepatitis C virus protease
JOURNAL Patent: US 5585259-A 85 17-DEC-1996;
FEATURES Location/Qualifiers
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/mol_type="unassigned DNA"

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QY 292 TGCACCTTCGGCTCCTCGGACCTTACCTGTGTACAGGACGCGCGATGTCATTC 347
Db 1000 TGCACCTTCGGCTCCTCGGACCTTACCTGTGTACAGGACGCGCGATGTCATTC 1055

RESULT 47
I34286 2523 bp DNA linear PAT 06-FEB-1997
LOCUS Sequence 85 from patent US 5597691.
DEFINITION I34286
ACCESSION I34286
VERSION I34286.1 GI:1825077
KEYWORDS SOURCE
ORGANISM Unknown.

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Unclassified.
1 (bases 1 to 2523)
Houghton, M., Choo, Q.-L. and Kuo, G.
Hepatitis C virus protease
Patent: US 5597691-A 85 28-JAN-1997;
Location/Qualifiers
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/mol_type="unassigned DNA"

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RESULT 48
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LOCUS      Definition
Accession      I82491
Version      I82491.1 GI:3210788
Keywords
Source
Organism      Unknown.
Unclassified.
REFERENCE      1 (bases 1 to 2523)
Authors      Houghton, M., Choo, Q.-L. and Kuo, G.
Title      Hepatitis C virus protease
Journal      Patent: US 5712145-A 85 27-JAN-1998;
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/mol_type="unassigned DNA"

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Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      292 TGCACCTTGGCGCTCCTCGGACCTTTACCTGGTCACGAGGACGCCGATGTCATTCC 347
Db      1000 TGCACCTTGGCGCTCCTCGGACCTTTACCTGGTCACGAGGACGCCGATGTCATTCC 1055

RESULT 49
AR118686      AR118686      5360 bp      DNA      linear      PAT 16-MAY-2001
LOCUS      Definition
Accession      AR118686
Version      AR118686.1 GI:14100596
Keywords
Source
Organism      Unknown.
Unclassified.
REFERENCE      1 (bases 1 to 5360)
Authors      Chien, D.Y.
Title      NANBV diagnostics and vaccines
Journal      Patent: US 6150087-A 53 21-NOV-2000;
Features      Location/Qualifiers
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RESULT 50
I06434      I06434      5360 bp      DNA      linear      PAT 02-DEC-1994
LOCUS      Definition
Accession      I06434
Version      I06434.1 GI:590311
Keywords
Source
Organism      Unknown.
Unclassified.
REFERENCE      1 (bases 1 to 5360)
Authors      Houghton, M., Choo, Q.-L. and Kuo, G.
Title      Nanbv diagnostics and vaccines
Journal      Patent: EP 0318216-A1 48 31-MAY-1989;
Features      Location/Qualifiers
source      1. .5360
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/mol_type="unassigned DNA"

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QY      292 TGCACCTTGGCGCTCCTCGGACCTTTACCTGGTCACGAGGACGCCGATGTCATTCC 347
Db      1221 TGCACCTTGGCGCTCCTCGGACCTTTACCTGGTCACGAGGACGCCGATGTCATTCC 1276

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Job time : 7832 secs
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GenCore version 5.1.6  
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OM nucleic - nucleic search, using sw model

Run on: August 19, 2004, 03:08:25 ; Search time 782 Seconds  
(without alignments)

11196.338 Million cell updates/sec

Title: US-09-930-591-1

Perfect score: 2061

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Gapop 60.0 , Gapext 60.0

Searched: 3373863 seqs, 2124099041 residues

Word size : 35

Total number of hits satisfying chosen parameters: 209

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 1000 summaries

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- 2: Geneseq1990s:\*\*
- 3: Geneseq2000s:\*\*
- 4: Geneseq2001as:\*\*
- 5: Geneseq2001bs:\*\*
- 6: Geneseq2002s:\*\*
- 7: Geneseq2003as:\*\*
- 8: Geneseq2003bs:\*\*
- 9: Geneseq2003cs:\*\*
- 10: Geneseq2004s:\*\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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1	2061	100.0	2061	6	Aad34500 Hepatitis
2	2061	100.0	2061	6	Aad31767 Hepatitis
3	2061	100.0	2061	9	Aad60868 Hepatitis
4	62	3.0	943	2	Aaq32984 HCV NS3/N
5	62	3.0	943	2	Aax84003 HCV E1
6	62	3.0	943	2	Aax16760 Hepatitis
7	62	3.0	7983	9	Aad93727 Hepatitis
8	62	3.0	9365	6	Aad25518 Hepatitis
9	61	3.0	957	3	Aaz36164 Nucleotid
10	58	2.8	382	2	Aaq38233 HCV-108
11	58	2.8	382	5	Aaf32234 HCV recom
12	58	2.8	1414	2	Aaq38232
13	58	2.8	1414	5	Aaf32233 HCV recom
14	58	2.8	1420	2	Aaq38234 Clone pHc
15	58	2.8	1420	5	Aaf32235 HCV recom
16	56	2.7	281	2	Aaq14299 Hepatitis
17	56	2.7	281	2	Aaq14361 Clone C8h
18	56	2.7	281	2	Aat59255 HCV prote
19	56	2.7	281	2	Aav04988 Nucleotid
20	56	2.7	281	2	Aax26393 Nucleotid
21	56	2.7	281	8	Acd44791 Hepatitis
22	56	2.7	281	8	Ada07864 Hepatitis
23	56	2.7	282	1	Aan90317 Hepatitis

24	56	2.7	282	2	AAQ80170	Hepatitis
25	56	2.7	283	1	AAQ92087	Sequence
26	56	2.7	368	2	AAQ14297	Hepatitis
27	56	2.7	368	2	AAQ14359	Clone C20
28	56	2.7	368	2	AAQ80168	Hepatitis
29	56	2.7	368	2	AAT59254	Hepatitis
30	56	2.7	368	2	AAV04986	Nucleotid
31	56	2.7	368	2	AAQ26391	Nucleotid
32	56	2.7	368	8	ACD44789	Hepatitis
33	56	2.7	368	8	ADA07860	Hepatitis
34	56	2.7	612	8	ABX15706	Anti-vira
35	56	2.7	1947	2	AAQ14304	Vector cf
36	56	2.7	2058	6	ABX15344	Hepatitis
37	56	2.7	2058	6	AAD29795	HCV-1 NS3
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53	56	2.7	2523	8	ACD44796	Hepatitis
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58	56	2.7	6905	1	AAN92103	Combined
59	56	2.7	7310	1	AAN92106	Composite
60	56	2.7	7310	1	AAN90336	Hepatitis
61	56	2.7	8316	2	AAQ98221	Hepatitis
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64	56	2.7	9133	2	AAZ07656	Nucleotid
65	56	2.7	9185	2	AAQ05956	Sense str
66	56	2.7	9185	2	AAQ10566	Hepatitis
67	56	2.7	9185	2	AAQ00459	Hepatitis
68	56	2.7	9185	2	AAQ26737	Hepatitis
69	56	2.7	9185	3	AAV75297	Sense str
70	56	2.7	9400	2	AAQ21744	Compiled
71	56	2.7	9401	2	AAT12710	Hepatitis
72	56	2.7	9401	2	AAT99981	HCV poly
73	56	2.7	9401	6	AAV09989	HCV poly
74	56	2.7	9401	6	AD355043	Hepatitis
75	53	2.6	475	6	ABL55592	HCV bait
76	53	2.6	583	6	ABL55594	HCV bait
77	53	2.6	790	6	ABL55593	HCV bait
78	53	2.6	836	6	ABL55591	HCV bait
79	53	2.6	7989	9	ADD93722	Hepatitis
80	53	2.6	7992	9	ADD93723	Hepatitis
81	53	2.6	9416	2	AAQ23871	NANRV Rut
82	53	2.6	9416	6	AAQ31764	Hepatitis
83	53	2.6	9416	9	AAD60865	Hepatitis
84	53	2.6	9518	5	AAQ03778	Hepatitis
85	53	2.6	9518	5	AAQ03808	Hepatitis
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87	53	2.6	9599	2	AAQ24832	Infectiou
88	53	2.6	9599	4	AAQ23491	Infectiou
89	53	2.6	9599	4	AAQ23491	Infectiou
90	53	2.6	9599	4	AAQ23491	Infectiou
91	53	2.6	9611	5	AAC86938	Nucleotid
92	53	2.6	9611	5	AAC86646	Nucleotid
93	53	2.6	9611	5	AAC86646	Nucleotid
94	53	2.6	9611	5	AAC86647	Nucleotid
95	53	2.6	9646	2	AAV59361	Nucleotid
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Qy	1801	TATAGACTGGGCGCTGTCCAGAATGAAGTCACCTGACGCAACCCAGTCACCAAGTATATC	1860
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Qy	1861	ATGACATGTATGTCGGCTGACCTGGAGGTGCTCACAGATACCTGGGTGCTCGTTGGCGGC	1920
Db	1861	ATGACATGTATGTCGGCTGACCTGGAGGTGCTCACAGATACCTGGGTGCTCGTTGGCGGC	1920
Qy	1921	GTTCCTGGCTCCTTTGGSCGGGTATGTCCTATCCACAGGCTCGCTGTGCATAGTAGTAGG	1980
Db	1921	GTTCCTGGCTCCTTTGGSCGGGTATGTCCTATCCACAGGCTCGCTGTGCATAGTAGTAGG	1980
Qy	1981	ATTGTCCTTGTCCGAAAGCCGGCAATCATACCGACAGGGAAGTCTCTACCCGGAGTTC	2040
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Qy	2041	GATGAAATGGAAGTAGTCTGA	2061
Db	2041	GATGAAATGGAAGTAGTCTGA	2061

RESULT 2	
AAD31767	
ID	AAD31767 standard; DNA; 2061 BP.
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XX	
AC	AAD31767;
XX	
DT	18-JUN-2002 (first entry)
XX	
DE	Hepatitis C virus (HCV) NS3/4A DNA coding region.
XX	
KW	Hepatitis C virus; HCV infection; virucide; fungicide; antibacterial;
KW	cytostatic; immunostimulant; vaccine; ribavirin; immune response; cancer;
KW	ds.
XX	
OS	Hepatitis C virus.
XX	
XX	
XX	
Key	Location/Qualifiers
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FT	/*tag= a
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XX	WO200213855-A2.
PN	
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XX	21-FEB-2002.
PD	
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PF	15-AUG-2001; 2001WO-IB001808.
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XX	17-AUG-2000; 2000US-0225767P.
PR	29-AUG-2000; 2000US-0229175P.
PR	03-NOV-2000; 2000US-00705547.
XX	
XX	(TRIP-) TRIPEP AB.
PA	
XX	
PI	Sallberg M, Hultgren C;
XX	
DR	WPI; 2002-241837/29.
DR	P-PSDB; AAE19900.
DR	
XX	
PT	Vaccine compositions for treating and preventing disease, preferably
PT	Hepatitis C virus infection, comprises ribavirin and antigen that has
PT	epitope present in hepatitis C virus.
XX	
XX	Claim 1; Page 94-95; 120pp; English.
PS	
XX	
CC	The invention relates to a composition comprising ribavirin and an
CC	antigen preferably non structural 3 protein (NS3)/4A fragment of
CC	hepatitis C virus (HCV) genome or a peptide or nucleic acid of HCV
CC	sequence. The composition is useful for enhancing an immune response to a
CC	hepatitis C antigen in humans, domestic, sport or pet species and as
CC	vaccines for treating and preventing HCV infections. The composition is

QY 961 CTGCGCACCAGTACCCCTCCGGGCTCCGTCACCTGTGCCCCCATCTTAACATCGAGGAGTT 1020  
 Db 961 CTGCGCACCAGTACCCCTCCGGGCTCCGTCACCTGTGCCCCCATCTTAACATCGAGGAGTT 1020  
 QY 1021 GCTCTGTCCACTACCGAGAGATCCCTTTTATGGCAAGGCTATTCCTTTGAAGCAAT 1080  
 Db 1021 GCTCTGTCCACTACCGAGAGATCCCTTTTATGGCAAGGCTATTCCTTTGAAGCAAT 1080  
 QY 1081 AAGGGGGGAGACATCTCTTCTGCGCACTCAAAAGAAAGTGCAGAGCTCGCGCA 1140  
 Db 1081 AAGGGGGGAGACATCTCTTCTGCGCACTCAAAAGAAAGTGCAGAGCTCGCGCA 1140  
 QY 1141 AACTGTGCTGCTGCGGCTCAATGCGGCTTACTACCGCGCTTGATGTCGCTC 1200  
 Db 1141 AACTGTGCTGCTGCGGCTCAATGCGGCTTACTACCGCGCTTGATGTCGCTC 1200  
 QY 1201 ATCCCGACCAAGTGTGCTGCTGCTGCGCACTGACGCCCTCATGACCGCTTTACC 1260  
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 Db 1321 CTTGACCCCTACCTTCACATTTGAGACATCAAGCTTCCCGAGATGCTGTCTCCGTA 1380  
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 Db 1381 CAACGTCGGGTAGACTGCGAGAGGAAGCCAGGCAATACAGATTTGTGGACCGGG 1440  
 QY 1441 GAGGCTCCTTCTGCGATTTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 1500  
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 QY 1681 CTGCTAGCTACCAAGCCACCGTGTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 1740  
 Db 1681 CTGCTAGCTACCAAGCCACCGTGTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 1740  
 QY 1741 CAGATGTGAAGTCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 1800  
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 QY 1801 TATAGACTGGGCGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 1860  
 Db 1801 TATAGACTGGGCGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 1860  
 QY 1861 ATGACATGTATGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 1920  
 Db 1861 ATGACATGTATGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 1920  
 QY 1921 GTTCTGGCTGCTTGGCGGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 1980  
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 QY 1981 ATTCTTGTTCGGAAGCGCGCAATCATACCGGACAGGGAAGTCTTCTTACCGGAGTTC 2040  
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Db 2041 GATGAATGGAAGAGTCTCTGA 2061  
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 ID AAD60868 standard; DNA; 2061 BP.  
 XX AAD60868;  
 AC AC  
 XX XX  
 DT 15-JAN-2004 (first entry)  
 XX  
 DE Hepatitis C virus NS3/4A DNA.  
 XX Ribavirin; vaccine; immune response; infection; therapy; immunostimulant;  
 KW virucide; ds.  
 XX  
 OS Hepatitis C virus.  
 XX  
 FH Key Location/Qualifiers  
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 FT /\*tag= a  
 FT /product= "Hepatitis C virus protein"  
 XX US2002136740-A1.  
 PN 26-SEP-2002.  
 XX  
 XX 15-AUG-2001; 2001US-00929955.  
 XX 17-AUG-2000; 2000US-0225767P.  
 PR 29-AUG-2000; 2000US-0229175P.  
 XX (SALL/) SALLBERG M.  
 PA (HULT/) HULTGREN C.  
 XX Sallberg M, Hultgren C;  
 DR WPI: 2003-764978/72.  
 DR P-PSDB; ABW00351.  
 XX Vaccine compositions for treating and preventing disease, preferably  
 PT hepatitis C virus infection, comprises ribavirin and antigen that has  
 PT epitope present in hepatitis C virus.  
 XX Claim 1: Page 60-61; Opp; English.  
 XX The invention relates to a composition comprising ribavirin and an  
 CC antigen, where the antigen is derived from a hepatitis virus. The vaccine  
 CC is useful in enhancing the immune response to a hepatitis C antigen where  
 CC the composition is delivered to an animal identified as requiring an  
 CC enhanced immune response. The vaccine is useful in the treatment and  
 CC prevention of hepatitis C infection. The present sequence is Hepatitis C  
 CC virus NS3/4A DNA  
 SQ Sequence 2061 BP; 427 A; 616 C; 571 G; 447 T; 0 U; 0 Other;  
 Query Match 100.0%; Score 2061; DB 9; Length 2061;  
 Best Local Similarity 100.0%; Pred. No. 0;  
 Matches 2061; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 ATGGCGCCTATCACGGCCTATGCCAGACAGACAGGGGCTTTTGGGATGCATATCACC 60  
 Db 1 ATGGCGCCTATCACGGCCTATGCCAGACAGACAGGGGCTTTTGGGATGCATATCACC 60  
 QY 61 ACCTTGACCGCGCGGACAAACACAGGTGGAGGTTCAGATCGTGTCAACTGCT 120  
 Db 61 ACCTTGACCGCGCGGACAAACACAGGTGGAGGTTCAGATCGTGTCAACTGCT 120  
 QY 121 GCCCAGACTTTCTTGGCAACTGCAATTAACGGGGTGTGTGGACTGTCTACCATGGAGCC 180  
 Db 121 GCCCAGACTTTCTTGGCAACTGCAATTAACGGGGTGTGTGGACTGTCTACCATGGAGCC 180

181 GGAAACAGGACCAATTGGTCACTTAAGGTCCTGTTATCCAGATGTACCAATGTGGAC 240  
181 GGAAACAGGACCAATTGGTCACTTAAGGTCCTGTTATCCAGATGTACCAATGTGGAC 240  
241 CAGACCTCGTAGGCTGGCGGTCCCAAGGTGCGGCTCATTAACACCATGCATTCG 300  
241 CAGACCTCGTAGGCTGGCGGTCCCAAGGTGCGGCTCATTAACACCATGCATTCG 300  
301 GGCTTCCTCGACCTTTACCTGGTCAAGAGCAGCGCATGTCATTCCTGTGCGCCGACG 360  
301 GGCTTCCTCGACCTTTACCTGGTCAAGAGCAGCGCATGTCATTCCTGTGCGCCGACG 360  
361 GGTGATGGCAGGCGAGCTGCTTTCCCGCGGCTATCTCTTACTTGAAGGCTCTCTG 420  
361 GGTGATGGCAGGCGAGCTGCTTTCCCGCGGCTATCTCTTACTTGAAGGCTCTCTG 420  
421 GGAGGCCCTCTGCTGTGCCCCGAGGACATGCCGTAGGCATATTCAAGAGCGCGGTATGC 480  
421 GGAGGCCCTCTGCTGTGCCCCGAGGACATGCCGTAGGCATATTCAAGAGCGCGGTATGC 480  
481 ACCCGTGGAGTGGTAAAGCGGTGGACTTCAATCCCGTAGAGAGCTTAGAGCAACCATG 540  
481 ACCCGTGGAGTGGTAAAGCGGTGGACTTCAATCCCGTAGAGAGCTTAGAGCAACCATG 540  
541 AGGTCCCGGTGTTCTCAGACAACTCTCCCAAGCAGCAGTGCACAGAGCTTACCAAGTG 600  
541 AGGTCCCGGTGTTCTCAGACAACTCTCCCAAGCAGCAGTGCACAGAGCTTACCAAGTG 600  
601 GCCCACCCTGATGCTCCCAAGCGGAGGTAAGAGCAACCAAGTTCGCGCGCATACGCA 660  
601 GCCCACCCTGATGCTCCCAAGCGGAGGTAAGAGCAACCAAGTTCGCGCGCATACGCA 660  
661 GCTCAGGCTACAGAGTGTGCTGCTCAACCCCTCGTTCGCTGCAACATGGGCTTTGGT 720  
661 GCTCAGGCTACAGAGTGTGCTGCTCAACCCCTCGTTCGCTGCAACATGGGCTTTGGT 720  
721 GCTTACATGTCCAAGGCCCATGGATTGATCTCAACATCAGGAGTGGGGTGAGGCAAAAT 780  
721 GCTTACATGTCCAAGGCCCATGGATTGATCTCAACATCAGGAGTGGGGTGAGGCAAAAT 780  
781 ACTACTGGCAGCCGATCAGATATCCACTACGGCAAGTTCCTGCGCAGCGCGGTGT 840  
781 ACTACTGGCAGCCGATCAGATATCCACTACGGCAAGTTCCTGCGCAGCGCGGTGT 840  
841 TCAGGGGTGCTTATGACATAAATAATTGTGACAGTGCCACTCCACGGATGCAACATCC 900  
841 TCAGGGGTGCTTATGACATAAATAATTGTGACAGTGCCACTCCACGGATGCAACATCC 900  
901 ATCTTGGGCATTTGGCACTGTCTTGACCAAGAGAGACCGCGGGGCGAGACTGACTGTG 960  
901 ATCTTGGGCATTTGGCACTGTCTTGACCAAGAGAGACCGCGGGGCGAGACTGACTGTG 960  
961 CTCGCCACCGCTACCCCTCGGGCTCGTCACTGTGCCCATCTTAACATCAGGAGGTT 1020  
961 CTCGCCACCGCTACCCCTCGGGCTCGTCACTGTGCCCATCTTAACATCAGGAGGTT 1020  
1021 GCTCTGTCCACTACCGGAGAGATCCCTTTTATGGCAAGGCTATTCCTTTGAAGCAAT 1080  
1021 GCTCTGTCCACTACCGGAGAGATCCCTTTTATGGCAAGGCTATTCCTTTGAAGCAAT 1080  
1081 AAGGGGGGAGACATCTCACTTCTGCACTCAAGAGAGTGGCAGAGCTCGCGCA 1140  
1081 AAGGGGGGAGACATCTCACTTCTGCACTCAAGAGAGTGGCAGAGCTCGCGCA 1140  
1141 AAACCTGTGCGGTTGGCGGTCAATGCGGTGCTTACTACCGCGGCTTGATGTCCGTC 1200  
1141 AAACCTGTGCGGTTGGCGGTCAATGCGGTGCTTACTACCGCGGCTTGATGTCCGTC 1200  
1201 ATCCCGACCAAGTGTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTTACC 1260  
1201 ATCCCGACCAAGTGTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTTACC 1260  
1261 GCGGACTTCGATTCGGTGTAGTAGTGCAACACGCTGTGTACCCAGACAGTCGACTTCAGC 1320

1261 GCGGACTTCGATTCGGTGTAGTAGTGCAACACGCTGTGTACCCAGACAGTCGACTTCAGC 1320  
1321 CTTGACCTTCACTTCCACCATAGACATACGCTTCCCAAGGATGCTGTCTCCGCTACT 1380  
1321 CTTGACCTTCACTTCCACCATAGACATACGCTTCCCAAGGATGCTGTCTCCGCTACT 1380  
1381 CAACTCGGGGTAGGACTGGCAGAGGAGCCAGCATCTACAGATTGTGGCACCAGGG 1440  
1381 CAACTCGGGGTAGGACTGGCAGAGGAGCCAGCATCTACAGATTGTGGCACCAGGG 1440  
1441 GAGCGTCTCTTGGCATGTTTGAATCTGCTCTCTGCGAGTGTATGACCGGGTGT 1500  
1441 GAGCGTCTCTTGGCATGTTTGAATCTGCTCTCTGCGAGTGTATGACCGGGTGT 1500  
1501 GCTTGGTATGAGCTTACGCCCGCAGACACAGATTAGGCTACGAGCATACATGAACACC 1560  
1501 GCTTGGTATGAGCTTACGCCCGCAGACACAGATTAGGCTACGAGCATACATGAACACC 1560  
1561 CCGGACTTCCCGTGTGCCAAGACCACTTCCATCCAGACAAAGCAGAGTGGGAAACCTTCCCTAT 1620  
1561 CCGGACTTCCCGTGTGCCAAGACCACTTCCATCCAGACAAAGCAGAGTGGGAAACCTTCCCTAT 1620  
1621 ACCCAGATAGACGCCCACTTCCATCCAGACAAAGCAGAGTGGGAAACCTTCCCTAT 1680  
1621 ACCCAGATAGACGCCCACTTCCATCCAGACAAAGCAGAGTGGGAAACCTTCCCTAT 1680  
1681 CTGGTAGGCTTACCAAGCCACCGTGTGCGCTAGAGCTCAAGCCCTCCCGCTGCTGGGAC 1740  
1681 CTGGTAGGCTTACCAAGCCACCGTGTGCGCTAGAGCTCAAGCCCTCCCGCTGCTGGGAC 1740  
1741 CAGATGTGGAAGTGTTCATCCGCTCTCAAGCCCACTTCCATGGGCGCAACACCTTCTGCTA 1800  
1741 CAGATGTGGAAGTGTTCATCCGCTCTCAAGCCCACTTCCATGGGCGCAACACCTTCTGCTA 1800  
1801 TATAGACTGGGCGTGTCCAGATGAAGTACCTGACGCAACCCAGTCAACCAAGTATATC 1860  
1801 TATAGACTGGGCGTGTCCAGATGAAGTACCTGACGCAACCCAGTCAACCAAGTATATC 1860  
1861 ATGACATGTATGCGGTGACCTGAGTGTCTGAGTGTCTGAGTGTCTGTTGGCGGC 1920  
1861 ATGACATGTATGCGGTGACCTGAGTGTCTGAGTGTCTGAGTGTCTGTTGGCGGC 1920  
1921 GTTCTGGCTGTGTTGGCGGTATTCCTATCCACAGGCTCGGTGGTCAATAGTAGTAGG 1980  
1921 GTTCTGGCTGTGTTGGCGGTATTCCTATCCACAGGCTCGGTGGTCAATAGTAGTAGG 1980  
1981 ATTGTCTTTCGGAAGCGGCAATCATACCCAGAGGAGTCTCTTACCGGAGTTC 2040  
1981 ATTGTCTTTCGGAAGCGGCAATCATACCCAGAGGAGTCTCTTACCGGAGTTC 2040  
2041 GATGAATGGAAGTGTCTGA 2061  
2041 GATGAATGGAAGTGTCTGA 2061

## RESULT 4

AAQ32984

ID AAQ32984 standard; DNA; 943 BP.

XX

AC AAQ32984;

XX

DT 24-OCT-2003 (revised)

DT

DT 25-MAR-2003 (revised)

DT

DT 14-MAY-1993 (first entry)

XX

XX HCV NS3/NS4 non-structural region.

XX

XX PCR; amplification; prototype; HCV pr; ss.

XX

XX Hepatitis C virus; HCVEL.

XX

XX FH Key Location/Qualifiers

FT CDS 2. .1210  
PT /\*tag= a  
PI /label= NS3/NS4

WO9221759-A1.  
10-DEC-1992.

04-JUN-1992; 92WO-FR000501.  
06-JUN-1991; 91FR-00006882.  
(INSP ) INST PASTEUR.

Brechet C, Kreamsdorf D, Porchon C;  
WPI; 1992-433657/52.

New nucleotide and peptide sequences - specific for French isolate of  
hepatitis C virus and useful in diagnosing and treating related  
infections.

Disclosure; Fig 8; 50pp; French.

RNA was extracted from the serum of an HCV-positive blood donor, subjected  
to reverse transcription and the cDNA formed amplified by PCR.  
Amplification prods. were cloned, screened with a probe derived  
from the HCV prototype and inserts sequenced. The results showed marked  
conservation in the non-coding region, significant variability in the  
structural region (encoding envelope proteins) and reduced variability in  
the non-structural region. The NS3/NS4 coding region corresponds to  
positions 4361-5303 of HCV prototype (HCV pt) (WO-A-90/14436). (Updated  
on 25-MAR-2003 to correct PN field.) (Updated on 24-OCT-2003 to  
standardise OS field)

Sequence 943 BP; 204 A; 292 C; 256 G; 191 T; 0 U; 0 Other;

Query Match 3.0%; Score 62; DB 2; Length 943;  
Best Local Similarity 100.0%; Pred. No. 8.5e-21;  
Matches 62; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1915 GCGCGCTTCTGGCTTTGGCGCGTATTGCTATCCACAGCGTGGTGGTCATAGTA 1974  
|||||  
630 GCGCGCTTCTGGCTTTGGCGCGTATTGCTATCCACAGCGTGGTGGTCATAGTA 689

1975 GG 1976  
||  
690 GG 691

RESULT 5  
AA84003  
ID AAX84003 standard; cDNA; 943 BP.

AC AAX84003;  
XX  
DT 27-AUG-2003 (revised)  
DT 26-AUG-1999 (first entry)  
XX  
DE HCV E1 coding sequence.  
XX  
KW HCV E1 region; monoclonal antibody; diagnosis; HCV E1-specific antigen;  
KW ss.

Hepatitis C virus.  
US919454-A.  
06-JUL-1999.  
07-JUN-1995; 95US-00487231.  
18-MAR-1993; 93US-00965285.

5,879.904

XX (INSP ) INST PASTEUR.  
XX Porchon C, Brechet C, Kreamsdorf D;  
XX WPI; 1999-394595/33.  
DR P-PSDB; AAY22022.  
XX  
PT Nucleotides and peptides from hepatitis C virus isolate for detecting E1-  
PT specific antigens.  
XX  
PS Disclosure; Col 15-18; 45pp; English.  
XX  
CC This sequence encodes a hepatitis c virus (HCV) E1 region protein. The  
CC invention relates to human or murine monoclonal antibodies directed  
CC against a HCV E1 protein sequence. The monoclonal antibodies and their  
CC fragments are useful for the in vitro diagnosis of HCV E1-specific  
CC antigens. (Updated on 27-AUG-2003 to correct OS field.)  
XX  
SQ Sequence 943 BP; 204 A; 292 C; 256 G; 191 T; 0 U; 0 Other;

Query Match 3.0%; Score 62; DB 2; Length 943;  
Best Local Similarity 100.0%; Pred. No. 8.5e-21;  
Matches 62; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1915 GCGCGCTTCTGGCTTTGGCGCGTATTGCTATCCACAGCGTGGTGGTCATAGTA 1974  
|||||  
630 GCGCGCTTCTGGCTTTGGCGCGTATTGCTATCCACAGCGTGGTGGTCATAGTA 689

1975 GG 1976  
||  
690 GG 691

RESULT 6  
AA16760  
ID AAX16760 standard; cDNA to mRNA; 943 BP.

AC AAX16760;  
XX  
DT 27-APR-1999 (first entry)  
XX  
DE Hepatitis C virus NS3/NS4 region.  
XX  
KW E1 region; French Hepatitis C virus; HCV; immunogen; antibody; detection;  
KW immunoassay; ss.  
XX  
OS Hepatitis C virus.

Key Location/Qualifiers  
CDS 3..941  
/\*tag= a  
/product= "NS3/NS4 protein"  
/note= "no start or stop codons are given at the 5' or 3'  
ends of the sequence"

US9866139-A.  
02-FEB-1999.  
07-JUN-1995; 95US-00483695.  
18-MAR-1993; 93US-00965285.  
(INSP ) INST PASTEUR.  
Porchon C, Kreamsdorf D, Brechet C;  
WPI; 1999-141865/12.  
P-PSDB; AAW75483.  
New isolated and purified Hepatitis C virus E1 peptides - useful for  
vaccine production or diagnostic purposes.

XX Disclosure; Col 15-18; 45pp; English.

PS The sequence represents the NS3/NS4 region from a French Hepatitis C

CC virus (HCV) isolate. The encoded protein or peptides derived from it can

CC be: (i) conjugated to a carrier protein and used as immunogens for

CC eliciting protective antibodies; or (ii) labelled, and used as

CC immunoassay reagents for detecting antibodies specific for HCV E1

XX

SQ Sequence 943 BP; 204 A; 292 C; 256 G; 191 T; 0 U; 0 Other;

Query Match 3.0%; Score 62; DB 2; Length 943;

Best Local Similarity 100.0%; Pred. No. 8.5e-21;

Matches 62; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1915 GCGCGGTTCTGGTCTTTGCGCGGTATTGCTATCCAGGCTCGGTGGTCATAGTA 1974

DB 630 GCGCGGTTCTGGTCTTTGCGCGGTATTGCTATCCAGGCTCGGTGGTCATAGTA 689

QY 1975 GG 1976

DB 690 GG 691

RESULT 7

ADD93727

ID ADD93727 standard; DNA; 7993 BP.

XX

AC ADD93727;

XX

DT 29-JAN-2004 (first entry)

XX

DE Hepatitis C virus genotype 1a replicon.

XX

KW HCV; vaccine; virucide; ss.

XX

OS Hepatitis C virus; genotype 1a.

XX

PN WO2003085084-A2.

XX

PD 16-OCT-2003.

XX

PF 03-APR-2003; 2003WO-US010177.

XX

PR 03-APR-2002; 2002US-0369685P.

XX

PA (SMIK ) SMITHLINE BEECHAM CORP.

XX

PI Gates A, Gu B, Sarisky RT;

XX

DR WPI; 2003-804301/75.

XX

PT New hepatitis C virus (HCV) sub-genomic replicon, useful for facilitating

PT screening or testing of anti-HCV drugs, comprises a nucleic acid

PT construct encoding chimeric HCV non-structural proteins, and an NS5B

PT polymerase gene.

XX

PS Claim 16; Page 51-56; 159pp; English.

XX

CC The present sequence comprises a replicating hepatitis C virus (HCV)

CC genotype 1a replicon. The invention provides sub-genomic replicons of HCV

CC comprising a nucleic acid construct encoding chimeric HCV nonstructural

CC protein and an NS5B polymerase gene. A preferred replicon comprises an

CC NS3 nucleotide sequence ADD93721 that encodes the first 75 contiguous N-

CC terminal amino acids of the NS3 of genotype 1b, of a BB7 strain. A

CC chimeric replicon may comprise an NS3 sequence from any of the 6 major

CC HCV genotypes and subtypes but has its first 225 nucleotides of the

CC coding sequence replaced by the BB7 strain NS3 sequence, especially where

CC the replicon is from HCV genotype 1a (H77 strain) or genotype 1b (J4

CC strain). Stable cell lines expressing and replicating functional

CC replicons containing sequences from HCV genotype 1a (strain H77) or

CC genotype 1b (strain J4) within the prototype 1b replicon backbone from

CC HCV strain BB7 are provided. These can be used to screen for compounds

CC that modulate viral replication. The sub-genomic HCV replicon systems of

CC the invention may provide the foundation for generating HCV replicons of

CC all 6 major genotypes and subtypes to facilitate screening, testing and

CC evaluating anti-infective agents for HCV disease(s).

XX

SQ Sequence 7993 BP; 1630 A; 2385 C; 2235 G; 1733 T; 0 U; 0 Other;

Query Match 3.0%; Score 62; DB 9; Length 7993;

Best Local Similarity 100.0%; Pred. No. 8.1e-21;

Matches 62; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 970 GTATCCCTCCGGCTCGTCACTGTGCCCATCTCTAATCATCGAGGTTGCTCTGTCC 1029

DB 2770 GTATCCCTCCGGCTCGTCACTGTGCCCATCTCTAATCATCGAGGTTGCTCTGTCC 2829

QY 1030 AC 1031

DB 2830 AC 2831

RESULT 8

AAD25518

ID AAD25518 standard; DNA; 9365 BP.

XX

AC AAD25518;

XX

DT 26-MAR-2002 (first entry)

XX

DE Hepatitis C virus isolate colonel complete DNA genome.

XX

KW HCV; hepatitis C virus; cytostatic; cancer; immunosuppressive; virucide;

KW antibacterial; fungicide; protozoicide; antirheumatic; antiinflammatory;

KW antiarthritic; rheumatoid arthritis; neuroprotective; multiple sclerosis;

KW immune response; vasotropic; vaccine; gene therapy; autoimmune disease;

KW vasculitis; ds.

XX

OS Hepatitis C virus.

XX

PN WO200176643-A1.

XX

PD 18-OCT-2001.

XX

PF 06-APR-2001; 2001WO-US011372.

XX

PR 07-APR-2000; 2000US-0195680P.

XX

PA (BAYU ) BAYLOR COLLEGE MEDICINE.

XX

PI Orson FM, Kinsey BM, Bhogal BS;

XX

DR WPI; 2002-066308/09.

XX

PT Composition for oral delivery of vaccines, comprises expression vector

PT containing antigenic genomic sequence, bound to aggregated protein-

PT polycationic polymer conjugate or suspension.

XX

PS Disclosure; Page 87-90; 145pp; English.

XX

CC The invention relates to a composition comprising an expression vector

CC bound to an aggregated protein-polycationic polymer conjugate or

CC suspension. The expression vector contains a promoter polynucleotide

CC sequence operatively linked to a polynucleotide sequence encoding an

CC antigen which is a fragment of a gene or genome associated with an

CC infectious disease, cancer and autoimmune disease such as rheumatoid

CC arthritis, vasculitis, and multiple sclerosis, pathogenic genomes

CC consisting of bacterium, fungus, protozoa and virus such as human

CC immunodeficiency virus (HIV), herpes simplex virus (HSV), hepatitis C

CC virus (HCV), influenza and respiratory syncytial virus (RSV), and

CC optionally comprising a nucleotide sequence encoding a cytokine (or a

CC cytokine expression vector), is useful for inducing an immune response

CC (systemic and/or mucosal) in an organism. The cytokine expression vector

CC contains a sequence for granulocyte macrophage-colony stimulating factor

CC (GM-CSF) or interleukin-12 (IL-12). The polynucleotide sequences encoding



CC the antigen and the cytokine are under transcriptional control of same or  
CC different promoter polynucleotide sequences. The expression vector, as a  
CC DNA vaccine is useful for treating a condition in an organism. The  
CC present sequence is hepatitis C virus isolate colonel complete DNA genome  
CC related to the invention

SQ Sequence 9365 BP; 1863 A; 2835 C; 2674 G; 1993 T; 0 U; 0 Other;

Query Match 3.0%; Score 62; DB 6; Length 9365;  
Best Local Similarity 100.0%; Pred. No. 8.1e-21;  
Matches 62; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 970 GCTACCCCTCGGCTCGTCACTGTCCCATCTTAACTCATGAGAGGTTGCTCTGTCC 1029  
DB 4374 GCTACCCCTCGGCTCGTCACTGTCCCATCTTAACTCATGAGAGGTTGCTCTGTCC 4433  
QY 1030 AC 1031  
DB 4434 AC 4435

RESULT 9  
AAZ36164  
ID AAZ36164 standard; DNA; 957 BP.  
AC AAZ36164;  
XX 11-FEB-2000 (first entry)  
DT Nucleotide sequence of the mtNFH6NS3 clone B9 fusion protein.  
DE HCV; NS3 helicase; HCV subtype 1a; HCV subtype 1b; HCV infection;  
KW solid phase immunoassay; HCV antigen; NS3 protease; HCV antibody;  
KW passive vaccination; ss.  
OS Synthetic.  
OS Hepatitis C virus.  
PH Key Location/Qualifiers  
FT 1. .957  
FT /\*tag= a  
FT /product= "mtNFH6NS3 clone B9 fusion protein"  
FT misc\_feature 1. .120  
FT /\*tag= b  
FT /note= "these nucleotides encode the non-NS3 sequence,  
FT which is the mtNF fusion partner, the hexanistidine tag  
FT and part of the multilinker"  
PN WO9954735-A1.  
XX 28-OCT-1999.  
PD 15-APR-1999; 99WO-BP002547.  
XX 17-APR-1998; 98EP-00870087.  
XX (INNO-) INNOGENETICS NV.  
XX Maertens G, Louwagie J, Bosman A, Sablon E, Zrein M;  
PI WPI; 2000-013283/01.  
DR P-PSDB; AAY43895.  
XX New hepatitis C-virus polypeptide used for treating the infection.  
PT Claim 22; Fig 3-1; 66pp; English.  
PS The present sequence encodes a fusion protein comprising a Hepatitis C  
CC virus (HCV) NS3 protein. The NS3 polypeptides are used in a solid phase  
CC immunoassay comprising a HCV antigen (preferably a NS3 helicase or NS3  
CC protease protein) in the presence of reducing agent on the solid phase.  
CC Use of the reducing agent and purification of the antigen using to HCV  
CC sulphonation and desulphonation steps increases its reactivity to HCV

CC antibodies and enables an earlier detection of HCV infection. The assay  
CC is used for detecting antibodies raised against the HCV antigen. The  
CC polypeptides are used for preventing and treating HCV infection. The  
CC polypeptides are also used for diagnosing hepatitis infection. The  
CC antibodies to these polypeptides are used for providing passive  
CC vaccination

SQ Sequence 957 BP; 211 A; 286 C; 259 G; 201 T; 0 U; 0 Other;

Query Match 3.0%; Score 61; DB 3; Length 957;  
Best Local Similarity 100.0%; Pred. No. 2.8e-20;  
Matches 61; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 710 TCGGCTTTGGTCTTACATGTCACAGGCCCATGGGATTCCTAACATCAGGACTGGG 769  
DB 344 TCGGCTTTGGTCTTACATGTCACAGGCCCATGGGATTCCTAACATCAGGACTGGG 403  
QY 770 T 770  
DB 404 T 404

RESULT 10  
AAQ38233  
ID AAQ38233 standard; DNA; 382 BP.  
XX AAQ38233;  
AC AAQ38233;  
XX 25-MAR-2003 (revised)  
DT 01-JUL-1993 (first entry)  
DT DE PHCV-108 coding for HCV amino acids 1454-1569.  
XX Hepatitis C virus; C100 antigen; CKS fusion protein; CMP-XDO synthetase;  
KW immunodot assay; Non-A, non-B hepatitis; PCR; polymerase chain reaction;  
KW ss.  
XX Hepatitis C virus.  
XX WO9304087-A1.  
PN 04-MAR-1993.  
PD 21-AUG-1992; 92WO-US007187.  
XX 21-AUG-1991; 91US-00748566.  
XX (ABBO ) ABBOTT LAB.  
XX Desai SM, Casey JM, Rupprecht KR, Devare SG;  
XX WPI; 1993-093940/11.  
XX Hepatitis C assay using recombinant C-100 region antigens - for detecting  
XX antibodies and antigen in body fluids from individuals exposed to  
XX hepatitis C virus.  
XX Example 11; Page 74; 206pp; English.  
XX Plasmid pHCV-68 (see AAQ38232) was used in the construction of pHCV-72  
CC (see AAR33569) which expresses the HCV CKS-200 antigen (HCV amino acids  
CC 1192-1931) at high levels in E. coli. An NcoI fragment containing the C100  
CC coding region was excised from plasmid pHCV-62 (i.e. a clone expressing  
CC the HCV CKS-C100 deletion antigen HCV amino acids 1569-1574 and 1598-  
CC 1931). This NcoI fragment was inserted into the NcoI site of pHCV-108  
CC (i.e. AAR38233, HCV amino acids 1454-1569) to create pHCV-68. The  
CC Clai/BamHI fragment of pHCV-68 containing the HCV NS3/C100 coding region  
CC was excised and inserted into the Clai/BamHI sites of pHCV-29 (i.e. a  
CC clone expressing HCV CKS-33C antigen). (Updated on 25-MAR-2003 to correct  
CC PN field.)  
XX Sequence 382 BP; 76 A; 113 C; 101 G; 92 T; 0 U; 0 Other;

Query Match 2.8%; Score 58; DB 2; Length 382;  
 Best Local Similarity 100.0%; Pred. No. 9.6e-19;  
 Matches 58; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1291 ACGTGTGTACCCAGACAGTCGACTTCAGCCTTGACCTTACCTTCACCATTTGAGACAA 1348  
 |||||  
 DB 33 ACGTGTGTACCCAGACAGTCGACTTCAGCCTTGACCTTACCTTCACCATTTGAGACAA 90  
 |||||

RESULT 11  
 AAF32233  
 ID AAF32234 standard; DNA; 382 BP.  
 AC AAF32234;  
 XX AAF32234;  
 DT 17-APR-2001 (first entry)  
 DE HCV recombinant antigen pHCV-108 amino acid sequence SEQ ID NO:56.  
 DE Hepatitis C virus; HCV; antigen; detection; antibody; ss.  
 OS Hepatitis C virus.  
 XX US6172189-B1.  
 PN US6172189-B1.  
 XX 09-JAN-2001.  
 XX 02-JUN-1997; 97US-00867611.  
 XX 24-AUG-1990; 90US-00572822.  
 XX 07-NOV-1990; 90US-00614069.  
 XX 21-AUG-1991; 91US-00748561.  
 XX 21-AUG-1991; 91US-00748565.  
 XX 21-AUG-1991; 91US-00748566.  
 XX 19-NOV-1992; 92US-00989843.  
 XX 10-JAN-1994; 94US-00179826.  
 XX 01-MAY-1996; 96US-00646757.  
 XX (ABBO) ABBOTT LAB.  
 XX Devare SG, Desai SM, Casey JM, Dailey SH, Dawson GJ;  
 PI Gutierrez RA, Lesniewski RR, Stewart JL, Rupprecht KR;  
 XX WPI; 2001-122352/13.  
 DR New recombinant antigens representing distinct antigenic regions of  
 XX Hepatitis C virus (HCV) genome, useful for detection of antibodies and  
 XX antigens in body fluids of individuals exposed to HCV.  
 FS Example 17; Col 203-204; 167pp; English.  
 XX The present invention describes recombinant Hepatitis C virus (HCV)  
 CC antigens (I). (I) is useful as a reagent for the detection of antibodies  
 CC and antigen in body fluids from individuals exposed to HCV. The HCV assay  
 CC uses reliable and efficient reagents and methods to accurately detect the  
 CC presence of HCV antibodies in samples obtained from individuals suspected  
 CC of having HCV infection. AAF32218 to AAF32235, AAB51371 to AAB51379 and  
 CC AAB69001 to AAB69032 represent sequences used in the exemplification of  
 CC the present invention  
 XX Sequence 382 BP; 76 A; 113 C; 101 G; 92 T; 0 U; 0 Other;

Query Match 2.8%; Score 58; DB 5; Length 382;  
 Best Local Similarity 100.0%; Pred. No. 9.6e-19;  
 Matches 58; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1291 ACGTGTGTACCCAGACAGTCGACTTCAGCCTTGACCTTACCTTCACCATTTGAGACAA 1348  
 |||||  
 DB 33 ACGTGTGTACCCAGACAGTCGACTTCAGCCTTGACCTTACCTTCACCATTTGAGACAA 90  
 |||||

RESULT 12  
 AAQ38232

ID AAQ38232 standard; DNA; 1414 BP.  
 XX AAQ38232;  
 AC AAQ38232;  
 DT 25-MAR-2003 (revised)  
 DT 01-JUL-1993 (first entry)  
 XX CKS-HCV antigen fusion gene pHCV-68.  
 DE Hepatitis C virus; C100 antigen; CKS fusion protein; CMP-KDO synthetase;  
 XX immunodot assay; Non-A, non-B hepatitis; ss.  
 KW Hepatitis C virus.  
 OS WO9304087-A1.  
 PN WO9304087-A1.  
 XX 04-MAR-1993.  
 PD 21-AUG-1992; 92WO-US007187.  
 XX 21-AUG-1991; 91US-00748566.  
 XX (ABBO) ABBOTT LAB.  
 XX Desai SM, Casey JM, Rupprecht KR, Devare SG;  
 PI WPI; 1993-093940/11.  
 XX Hepatitis C assay using recombinant C-100 region antigens - for detecting  
 PT antibodies and antigen in body fluids from individuals exposed to  
 PT hepatitis C virus.  
 XX Example 1; Page 61-62; 206pp; English.  
 XX Plasmid pHCV-68 was used in the construction of pHCV-72 (see AAR33569)  
 CC which expresses the HCV CKS-200 antigen (HCV amino acids 1192-1931) at  
 CC high levels in E.coli. An NcoI fragment containing the C100 coding region  
 CC was excised from plasmid pHCV-62 (i.e. a clone expressing the HCV CKS-  
 CC C100 deletion antigen HCV amino acids 1569-1574 and 1598-1931). This NcoI  
 CC fragment was inserted into the NcoI site of pHCV-108 (i.e. HCV amino  
 CC acids 1454-1569) to create pHCV-68. The ClaI/BamHI fragment of pHCV-68  
 CC containing the HCV NS3/C100 coding region was excised and inserted into  
 CC the ClaI/BamHI sites of pHCV-29 (i.e. a clone expressing HCV CKS-33C  
 CC antigen). (Updated on 25-MAR-2003 to correct PN field.)  
 XX Sequence 1414 BP; 244 A; 395 C; 387 G; 388 T; 0 U; 0 Other;

Query Match 2.8%; Score 58; DB 2; Length 1414;  
 Best Local Similarity 100.0%; Pred. No. 9.3e-19;  
 Matches 58; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1291 ACGTGTGTACCCAGACAGTCGACTTCAGCCTTGACCTTACCTTCACCATTTGAGACAA 1348  
 |||||  
 DB 22 ACGTGTGTACCCAGACAGTCGACTTCAGCCTTGACCTTACCTTCACCATTTGAGACAA 79  
 |||||

RESULT 13  
 AAF32233  
 ID AAF32233 standard; DNA; 1414 BP.  
 XX AAF32233;  
 AC AAF32233;  
 DT 17-APR-2001 (first entry)  
 DE HCV recombinant antigen pHCV-68 amino acid sequence SEQ ID NO:51.  
 DE Hepatitis C virus; HCV; antigen; detection; antibody; ss.  
 OS Hepatitis C virus.  
 XX US6172189-B1.  
 PN US6172189-B1.  
 XX 09-JAN-2001.

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XX PF 02-JUN-1997; 97US-00867611.
XX PR 24-AUG-1990; 90US-00572822.
XX PR 07-NOV-1990; 90US-00614069.
XX PR 21-AUG-1991; 91US-00748561.
XX PR 21-AUG-1991; 91US-00748565.
XX PR 21-AUG-1991; 91US-00748566.
XX PR 19-NOV-1992; 92US-00989843.
XX PR 10-JAN-1994; 94US-00179896.
XX PR 01-MAY-1996; 96US-00646757.
XX PA (ABBO ) ABBOTT LAB.
XX PI Devare SG, Desai SM, Casey JM, Dailey SH, Dawson GJ;
XX PI Gutierrez RA, Lesniewski RR, Stewart JL, Rupprecht KR;
XX DR WPI; 2001-122352/13.
XX
XX New recombinant antigens representing distinct antigenic regions of
XX PT Hepatitis C virus (HCV) genome, useful for detection of antibodies and
XX PT antigens in body fluids of individuals exposed to HCV.
XX PS Example 17; Col 185-188; 167pp; English.
XX
XX The present invention describes recombinant Hepatitis C virus (HCV)
XX CC antigens (I). (I) is useful as a reagent for the detection of antibodies
XX CC and antigen in body fluids from individuals exposed to HCV. The HCV assay
XX CC uses reliable and efficient reagents and methods to accurately detect the
XX CC presence of HCV antibodies in samples obtained from individuals suspected
XX CC of having HCV infection. AAF32235 to AAF32235, AAF51371 to AAF51379 and
XX CC AAB69001 to AAB69032 represent sequences used in the exemplification of
XX CC the present invention
XX
XX SQ Sequence 1414 BP; 244 A; 395 C; 387 G; 388 T; 0 U; 0 Other;
Query Match 2.8%; Score 58; DB 5; Length 1414;
Best Local Similarity 100.0%; Pred. No. 9.3e-19;
Matches 58; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1291 ACGTGTGTCACCCAGACAGTCGACTTCAGCCTTGACCTACCTACCTACCATTTGAGACAA 1348
Db 22 ACGTGTGTCACCCAGACAGTCGACTTCAGCCTTGACCTACCTACCTACCATTTGAGACAA 79
RESULT 14
AAQ38234
ID AAQ38234 standard; DNA; 1420 BP.
AC AAQ38234;
XX
XX 25-MAR-2003 (revised)
DT 01-JUL-1993 (first entry)
XX
XX Clone pHCV-69 containing the HCV NS3/C100 coding region.
XX
XX Hepatitis C virus; C100 antigen; CKS fusion protein; CMP-KDO synthetase;
XX KW immunodot assay; Non-A, non-B hepatitis; PCR; polymerase chain reaction;
XX KW ss.
XX
XX Hepatitis C virus.
XX OS
XX WO9304087-A1
XX
XX 04-MAR-1993.
XX
XX 21-AUG-1992; 92WO-US007187.
XX
XX 21-AUG-1991; 91US-00748566.
XX
XX (ABBO ) ABBOTT LAB.
XX
XX Desai SM, Casey JM, Rupprecht KR, Devare SG;

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XX WPI; 1993-093940/11.
XX
XX Hepatitis C assay using recombinant C-100 region antigens - for detecting
XX PT antibodies and antigen in body fluids from individuals exposed to
XX PT hepatitis C virus.
XX
XX Example 11; Page 74-75; 206pp; English.
XX
XX Plasmid pHCV-69 (i.e. AAQ38234) was used in the construction of pHCV-73
XX CC (see AAR3570) which expresses the HCV CKS-200 antigen (HCV amino acids
XX CC 1192-1599 and 1621-1931) at high levels in E.coli. The C100 coding region
XX CC was excised from plasmid pHCV-63 and inserted into pHCV-108 (i.e.
XX CC AAQ38233, HCV amino acids 1454-1569) to create pHCV-69. The ClaI/BamHI
XX CC fragment of pHCV-69 containing the HCV NS3/C100 coding region was excised
XX CC and inserted into the ClaI/BamHI sites of pHCV-29 (i.e. a clone
XX CC expressing HCV CKS-33C antigen). (Updated on 25-MAR-2003 to correct PN
XX CC field.)
XX
XX SQ Sequence 1420 BP; 246 A; 396 C; 384 G; 394 T; 0 U; 0 Other;
Query Match 2.8%; Score 58; DB 2; Length 1420;
Best Local Similarity 100.0%; Pred. No. 9.3e-19;
Matches 58; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1291 ACGTGTGTCACCCAGACAGTCGACTTCAGCCTTGACCTACCTACCTACCATTTGAGACAA 1348
Db 22 ACGTGTGTCACCCAGACAGTCGACTTCAGCCTTGACCTACCTACCTACCATTTGAGACAA 79
RESULT 15
AAF32235
ID AAF32235 standard; DNA; 1420 BP.
XX
XX AAF32235;
XX
XX 17-APR-2001 (first entry)
DT
XX
XX HCV recombinant antigen pHCV-59 amino acid sequence SEQ ID NO:57.
DE
XX
XX Hepatitis C virus; HCV; antigen; detection; antibody; ss.
XX
XX Hepatitis C virus.
XX
XX US6172189-B1.
XX
XX 09-JAN-2001.
XX
XX 02-JUN-1997; 97US-00857611.
XX
XX 24-AUG-1990; 90US-00572822.
XX PR 07-NOV-1990; 90US-00614069.
XX PR 21-AUG-1991; 91US-00748561.
XX PR 21-AUG-1991; 91US-00748565.
XX PR 21-AUG-1991; 91US-00748566.
XX PR 19-NOV-1992; 92US-00989843.
XX PR 10-JAN-1994; 94US-00179896.
XX PR 01-MAY-1996; 96US-00646757.
XX
XX (ABBO ) ABBOTT LAB.
XX
XX Devare SG, Desai SM, Casey JM, Dailey SH, Dawson GJ;
XX PI Gutierrez RA, Lesniewski RR, Stewart JL, Rupprecht KR;
XX
XX WPI; 2001-122352/13.
XX
XX New recombinant antigens representing distinct antigenic regions of
XX PT Hepatitis C virus (HCV) genome, useful for detection of antibodies and
XX PT antigens in body fluids of individuals exposed to HCV.
XX
XX Example 17; Col 203-204; 167pp; English.
XX
XX The present invention describes recombinant Hepatitis C virus (HCV)

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KW HCV; NS3; non-structural domain 3; protease; polyprotein; inhibitor;  
 XX screen; processing; infection; treatment; probe; hepatitis C virus; ss.  
 OS Hepatitis C virus; Virus.  
 XX US5585258-A.  
 XX 17-DEC-1996.  
 XX 06-DEC-1994; 94US-00350884.  
 XX 04-APR-1990; 90US-00505433.  
 PR 04-APR-1991; 91US-00680296.  
 XX (CHIR ) CHIRON CORP.  
 XX Choo Q, Kuo G, Houghton M;  
 XX WPI; 1997-051175/05.  
 DR P-PSDB; AAW01691.  
 XX Compsn. contg. hepatitis C virus NS3 domain protease and related fusion  
 PT proteins - useful for screening specific inhibitors, potential antiviral  
 PT agents, prepn. of antibodies and for cleaving specific poly(peptide)s).  
 XX Example 3; Col 65-68; 68pp; English.  
 XX Compsn. comprising the hepatitis C virus (HCV) NS3 domain protease or  
 CC its active truncation analogues are claimed. Also new are fusion proteins  
 CC comprising the protease (or analogues) and, e.g. human superoxide (SOD)  
 CC or ubiquitin. The protease is essential for polyprotein processing, and  
 CC thus infectivity, in HCV. The compsns. are used to screen for specific  
 CC inhibitors (possibly useful as antiviral agents), to generate specific  
 CC antibodies and to cleave specific polypeptides. HCV cDNA clones (AAV59250  
 CC - 56 encoding AAW01686-92 resp.) were isolated from HCV genomic library  
 CC using probes AAT59244-49. The clones were used in the preparation of full  
 CC length SOD-protease fusion proteins. (Updated on 25-MAR-2003 to correct  
 CC PP field.) (Updated on 17-OCT-2003 to standardise OS field)  
 XX SQ Sequence 281 BP; 41 A; 89 C; 92 G; 59 T; 0 U; 0 Other;  
 Query Match 2.7%; Score 56; DB 2; Length 281;  
 Best Local Similarity 100.0%; Pred. No. 1e-17;  
 Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 292 TGCACTTGGGCTCCTCGGACCTTTACCTGGTCAAGGACGCCGCGATGTCATTC 347  
 Db 4 TGCACCTTGGGCTCCTCGGACCTTTACCTGGTCAAGGACGCCGCGATGTCATTC 59  
 RESULT 19  
 AAV04988  
 ID AAV04988 standard; DNA; 281 BP.  
 XX AAV04988;  
 XX 27-AUG-2003 (revised)  
 DT 07-MAY-1998 (first entry)  
 XX Nucleotide sequence of the Hepatitis c virus (HCV) clone c8h.  
 DE  
 DE Protease; HCV; NS3 domain; human superoxide dismutase; fusion protein;  
 KW ubiquitin; assay; activity; anti-HCV; ss.  
 XX Hepatitis C virus.  
 XX Key Location/Qualifiers  
 FH 1. .279  
 FT CDS /\*tag= a  
 FT  
 XX US5712145-A.  
 XX 27-JAN-1998.

XX 06-SEP-1996; 96US-00709173.  
 XX 04-APR-1990; 90US-00505433.  
 PR 04-APR-1991; 91US-00680296.  
 PR 06-DEC-1994; 94US-00350884.  
 PR 12-MAY-1995; 95US-00440548.  
 XX (CHIR ) CHIRON CORP.  
 XX Choo Q, Kuo G, Houghton M;  
 XX WPI; 1998-119886/11.  
 DR P-PSDB; AAW46392.  
 XX Recombinant hepatitis C virus protease - useful in screening drugs for  
 PT activity against hepatitis C virus.  
 XX Disclosure; Fig 4; 68pp; English.  
 XX The present sequence represents the nucleotide sequence of the Hepatitis  
 CC C virus (HCV) clone c8h. The clone was isolated using hybridisation probe  
 CC AAV04975. A cDNA fragment encoding protease was isolated from the clone,  
 CC and cloned into an expression vector to produce a fusion protein with  
 CC human superoxide dismutase-protease. The HCV protease is believed to  
 CC cleave itself from the genomic polyprotein. In the absence of protease  
 CC activity, the HCV polyprotein should remain in its unprocessed form, and  
 CC thus render the virus non-infectious. Inhibitors of protease activity  
 CC should therefore also inhibit viral infectivity. The protease can  
 CC therefore be used for assaying compounds for activity against HCV.  
 CC (Updated on 27-AUG-2003 to correct OS field.)  
 XX SQ Sequence 281 BP; 41 A; 89 C; 92 G; 59 T; 0 U; 0 Other;  
 Query Match 2.7%; Score 56; DB 2; Length 281;  
 Best Local Similarity 100.0%; Pred. No. 1e-17;  
 Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 292 TGCACTTGGGCTCCTCGGACCTTTACCTGGTCAAGGACGCCGCGATGTCATTC 347  
 Db 4 TGCACCTTGGGCTCCTCGGACCTTTACCTGGTCAAGGACGCCGCGATGTCATTC 59  
 RESULT 20  
 AAX26393  
 ID AAX26393 standard; DNA; 281 BP.  
 XX AAX26393;  
 XX 26-MAY-1999 (first entry)  
 DT Nucleotide sequence of HCV protease clone c8h.  
 DE  
 DE HCV NS3 protease; truncation analog; HCV control; protease activity;  
 KW viral infectivity; inactive non-cleaving protease; ss.  
 XX Hepatitis C virus.  
 XX US5885799-A.  
 XX 23-MAR-1999.  
 XX 06-SEP-1996; 96US-00709177.  
 XX 04-APR-1990; 90US-00505433.  
 PR 04-APR-1991; 91US-00680296.  
 PR 06-DEC-1994; 94US-00350884.  
 PR 12-MAY-1995; 95US-00440548.  
 XX (CHIR ) CHIRON CORP.  
 XX Choo Q, Kuo G, Houghton M;  
 XX

Same as 9115575A

DR WPI: 1999-228536/19.  
 DR P-PSDB; AA97604.  
 XX  
 PT Preparation of new Hepatitis C Virus NS3 protease - useful for screening  
 PT for compounds which inhibit HCV infectivity.  
 XX  
 PS Example 3; Fig 4; 71pp; English.  
 XX  
 CC The specification describes a method for making a purified Hepatitis C  
 CC virus (HCV) NS3 protease or active truncation analog. If the HCV protease  
 CC N-terminal cleavage signal is excluded (so that self-cleavage is  
 CC prevented), the HCV protease remains in its unprocessed form, and renders  
 CC the virus noninfectious. The protease is therefore useful for assaying  
 CC pharmaceutical agents for control of HCV, as compounds which inhibit  
 CC protease activity sufficiently will also inhibit viral infectivity. An  
 CC inactive non-cleaving protease can be used to screen for inhibitors.  
 CC Recombinant expression systems can be utilised to prepare recombinant HCV  
 CC which can be used to produce monoclonal antibodies. The present sequence  
 CC was isolated in the course of the invention  
 XX  
 SQ Sequence 281 BP; 41 A; 89 C; 92 G; 59 T; 0 U; 0 Other;  
 Query Match 2.7%; Score 56; DB 2; Length 281;  
 Best Local Similarity 100.0%; Pred. NO. 1e-17;  
 Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 292 TGCACCTTGGCGCTCTCGGACCTTTACTGCTGTCACGAGGCACGCCGATGTCATTCC 347  
 DB 4 TGCACCTTGGCGCTCTCGGACCTTTACTGCTGTCACGAGGCACGCCGATGTCATTCC 59  
 RESULT 21  
 ACDA4791  
 ID ACD44791 standard; DNA; 281 BP.  
 XX  
 AC ACD44791;  
 XX  
 DT 09-SEP-2003 (first entry)  
 XX  
 DE Hepatitis C virus (HCV) protease clone C8h DNA.  
 XX  
 KW Hepatitis C virus; HCV; protease; protease inhibition; viral infection;  
 KW Gene; ds.  
 XX  
 OS Hepatitis C virus.  
 XX  
 FN US2003027317-A1.  
 XX  
 XX 06-FEB-2003.  
 PD  
 XX 18-JUN-2001; 2001US-00884456.  
 PF  
 XX 04-APR-1990; 90US-00505433.  
 PR 04-APR-1991; 91US-00680296.  
 PR 06-DEC-1994; 94US-00350884.  
 PR 12-MAY-1995; 95US-00440548.  
 PR 06-SEP-1996; 96US-00709177.  
 PR 19-FEB-1999; 99US-00253230.  
 XX  
 XX (HOUG/) HOUGHTON M.  
 PA (CHOO/) CHOO Q.  
 PA (KUOG/) KUO G.  
 XX  
 FI Houghton M, Choo Q, Kuo G;  
 XX  
 XX WPI: 2003-492037/58.  
 DR P-PSDB; ABC27015.  
 DR  
 XX New compositions comprising a hepatitis C virus (HCV) protease  
 PT polynucleotide, useful for assaying pharmaceutical agents for controlling  
 PT HCV, and as compounds which inhibit the protease activity and viral  
 PT infectivity.  
 XX

PS Example 3; Fig 4; 41pp; English.  
 XX  
 CC The invention describes a composition comprising a polynucleotide which  
 CC encodes only the hepatitis C virus (HCV) protease or an active HCV  
 CC protease analogue. The protease is useful for assaying pharmaceutical  
 CC agents for controlling HCV, and as compounds which inhibit the protease  
 CC activity sufficiently will also inhibit viral infectivity. This sequence  
 CC encodes hepatitis C virus (HCV) protease  
 XX  
 SQ Sequence 281 BP; 41 A; 89 C; 92 G; 59 T; 0 U; 0 Other;  
 Query Match 2.7%; Score 56; DB 8; Length 281;  
 Best Local Similarity 100.0%; Pred. NO. 1e-17;  
 Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 292 TGCACCTTGGCGCTCTCGGACCTTTACTGCTGTCACGAGGCACGCCGATGTCATTCC 347  
 DB 4 TGCACCTTGGCGCTCTCGGACCTTTACTGCTGTCACGAGGCACGCCGATGTCATTCC 59  
 RESULT 22  
 ADA07864  
 ID ADA07864 standard; cDNA; 281 BP.  
 XX  
 AC ADA07864;  
 XX  
 DT 06-NOV-2003 (first entry)  
 XX  
 DE Hepatitis C virus cDNA C8h encoding an NS3 protease fragment.  
 XX  
 KW ss; HCV; virucide; NS3 protease; serine protease; hSOD;  
 KW superoxide dismutase; yeast a-factor; interleukin-28; ubiquitin;  
 KW beta-galactosidase; beta-lactamase; horseradish peroxidase;  
 KW glucose oxidase; urease; HCV infection.  
 XX  
 OS Hepatitis C virus.  
 XX  
 PN US2003064499-A1.  
 XX  
 PD 03-APR-2003.  
 XX  
 PF 18-JUN-2001; 2001US-00884455.  
 XX  
 PR 04-APR-1990; 90US-00505433.  
 PR 04-APR-1991; 91US-00680296.  
 PR 06-DEC-1994; 94US-00350884.  
 PR 12-MAY-1995; 95US-00440548.  
 PR 06-SEP-1996; 96US-00709177.  
 PR 18-FEB-1999; 99US-00253675.  
 XX  
 XX (HOUG/) HOUGHTON M.  
 PA (CHOO/) CHOO Q.  
 PA (KUOG/) KUO G.  
 XX  
 FI Houghton M, Choo Q, Kuo G;  
 XX  
 XX WPI: 2003-540789/51.  
 DR P-PSDB; ADA07865.  
 DR  
 XX A composition for assaying and designing antiviral agents specific for  
 PT Hepatitis C virus (HCV) comprises a purified proteolytic polypeptide from  
 PT HCV or a polynucleotide which encodes HCV protease.  
 XX  
 PS Example 3; Fig 4; 40pp; English.  
 XX  
 CC The invention relates to a composition comprising a purified proteolytic  
 CC polypeptide derived from Hepatitis C virus (HCV) or a polynucleotide  
 CC which encodes only the HCV protease or an active HCV protease analogue,  
 CC or which encodes a fusion protein comprising HCV protease or HCV protease  
 CC analogue, and a fusion partner. Also included are a fusion protein  
 CC comprising a fusion partner fused to a proteolytic polypeptide derived  
 CC from HCV, a method for assaying compounds for activity against HCV  
 CC (comprising providing an active HCV protease, contacting the protease

CC with a compound capable of inhibiting serine protease activity and  
 CC measuring inhibition of the proteolytic activity of the HCV protease) and  
 CC an expression vector for producing HCV protease or HCV protease analogues  
 CC in a host cell (comprising a polynucleotide encoding HCV protease or an  
 CC HCV protease analogue, transcriptional and translational regulatory  
 CC sequences functional in the host cell operably linked to the HCV protease  
 CC -encoding polynucleotide and a selectable marker). The fusion partner is  
 CC selected from hSOD (human superoxide dismutase), yeast a-factor,  
 CC interleukin (IL)-2S, ubiquitin, beta-galactosidase, beta-lactamase,  
 CC horseradish peroxidase, Glucose oxidase and urease. The composition is  
 CC useful in assaying and designing antiviral agents specific for HCV. The  
 CC method is used in identifying antiviral agents effective for treating  
 CC HCV. The present sequence is a cDNA encoding an HCV NS3 protease or  
 CC fragment.

SQ Sequence 281 BP; 41 A; 89 C; 92 G; 59 T; 0 U; 0 Other;

Query Match 2.7%; Score 56; DB 8; Length 281;  
 Best Local Similarity 100.0%; Pred. No. 1e-17;  
 Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTGGCGCTCCTCGGACCTTTACCTGGTCACGAGGACGCGGATGTCATTC 347  
 Db 4 TGCACCTTGGCGCTCCTCGGACCTTTACCTGGTCACGAGGACGCGGATGTCATTC 59

RESULT 23

AA090317  
 ID AA090317 standard; DNA; 282 BP.

AC AA090317;

DT 25-MAR-2003 (revised)

DT 01-NOV-1989 (first entry)

XX Hepatitis C virus (HCV) cDNA in clone 8h.

XX Hepatitis C virus; cDNA; clone 8h; clone 33c; probe; vaccine.

XX Pan troglodytes.

XX Key Location/Qualifiers  
 CDS 3..281  
 /\*tag= a

FT misc\_feature 212..283

FT /\*tag= b

XX GB2212511-A.

XX 26-JUL-1989.

XX 18-NOV-1988; 88GB-00027024.

XX 18-NOV-1987; 87US-00122714.

XX 30-DEC-1987; 87US-00139886.

XX 26-FEB-1988; 88US-00161072.

XX 26-OCT-1988; 88US-00263584.

XX (CHIR ) CHIRON CORP.

XX Houghton M, Choo QL, Kuo G;

XX WPI; 1989-215054/30.

XX P-PSDB; AAP90148.

XX Hepatitis C virus gene - used for prodn. of polynucleotide probes

FT polypeptide(s) and antibodies for diagnosis, prevention and treatment of  
 PT infection.

XX Disclosure; Fig 16; 30pp; English.

XX The sequence shows hepatitis C virus (HCV) cDNA in clone 8h. The cDNA  
 CC encodes antigens which react with antibodies in patients with non-A non-B

CC hepatitis (NANBH). The cDNA can be used to design probes, or to  
 CC synthesise polypeptides, which are used to diagnose HCV-induced NANBH, to  
 CC raise antibodies for immunoassay or treatment, or to produce vaccines.  
 CC The misc. feature indicates overlap with clone 33c. See also AAP90148,  
 CC AA090303-15, and N903018-36. (Updated on 25-MAR-2003 to correct PD  
 CC field.) (Updated on 25-MAR-2003 to correct PR field.) (Updated on 25-MAR-  
 CC 2003 to correct PA field.)

SQ Sequence 282 BP; 41 A; 89 C; 92 G; 60 T; 0 U; 0 Other;

Query Match 2.7%; Score 56; DB 1; Length 282;  
 Best Local Similarity 100.0%; Pred. No. 1e-17;  
 Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTGGCGCTCCTCGGACCTTTACCTGGTCACGAGGACGCGGATGTCATTC 347  
 Db 6 TGCACCTTGGCGCTCCTCGGACCTTTACCTGGTCACGAGGACGCGGATGTCATTC 61

RESULT 24

AA080170

ID AA080170 standard; DNA; 282 BP.

AC AA080170;

XX 25-MAR-2003 (revised)

DT 17-AUG-1995 (first entry)

XX Hepatitis C virus (HCV) protease clone C8h DNA.

XX Hepatitis C virus protease; HCV; clone C8h; viral infectivity inhibition;  
 ds.

XX Hepatitis C virus.

XX Key Location/Qualifiers  
 FT mat\_peptide 1..280  
 FT /\*tag= a

XX US5371017-A

XX 06-DEC-1994.

XX 04-APR-1991; 91US-00680296.

XX 04-APR-1990; 90US-00505433.

XX (CHIR ) CHIRON CORP.

XX Houghton M, Choo Q, Kuo G;

XX WPI; 1995-021889/03.

XX P-PSDB; AAR68542.

XX DNA encoding hepatitis C virus protease - used to produce large amts. of  
 the protease and to develop prods. for inhibition of viral infectivity.

XX Example 3; Fig 4; 69pp; English.

XX AA080170 encodes AAR68542 hepatitis C virus (HCV) protease clone C8h,  
 CC using recombinant expression systems large amounts of protease can be  
 CC produced. The HCV protease can be used in the production of Abs. It can  
 CC also be used for assaying agents which inhibit protease activity, to  
 CC identify compounds which inhibit viral infectivity. (Updated on 25-MAR-  
 CC 2003 to correct PF field.)

SQ Sequence 282 BP; 41 A; 89 C; 92 G; 59 T; 0 U; 1 Other;

Query Match 2.7%; Score 56; DB 2; Length 282;  
 Best Local Similarity 100.0%; Pred. No. 1e-17;  
 Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTGGCGCTCCTCGGACCTTTACCTGGTCACGAGGACGCGGATGTCATTC 347





XX Hepatitis C virus; HCV; human superoxide dismutase; SOD; ds.

XX Hepatitis C virus.

XX WO9115596-A.

XX 17-OCT-1991.

XX 04-APR-1990; 90US-00505434.

XX 04-APR-1990; 90US-00505434.

XX (PROT-) PROTOS INC.

XX Rosenberg S;

XX WPI; 1991-325236/44.

XX P-PSDB; AAR14350.

XX Method for assaying pharmaceutical cpds. - for determining anti-Hepatitis

XX C Virus activity, using binding affinity.

XX Example 3; Fig 2; 68pp; English.

XX The DNA from the clone was used to prepare a hSOD:HCV protease fusion  
CC construct. The truncated protease analogue expressed by the resulting  
CC vector is proteolytically inactive and can be used to assay a wide range  
CC of pharmaceutical agents for controlling HCV. Those agents which inhibit  
CC the protease activity sufficiently will also inhibit viral infectivity.  
CC See also AAQ14358-Q14365

XX Sequence 368 BP; 70 A; 114 C; 108 G; 76 T; 0 U; 0 Other;

Query Match 2.7%; Score 56; DB 2; Length 368;

Best Local Similarity 100.0%; Pred. No. 1e-17;

Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTGGCGCTCCTCGGACCTTTACCTGTCACGAGGCACGCCGATGTCATTCC 347

Db 220 TGCACCTTGGCGCTCCTCGGACCTTTACCTGTCACGAGGCACGCCGATGTCATTCC 275

RESULT 28

AAQ80168

ID AAQ80168 standard; DNA; 368 BP.

XX AAQ80168;

XX 25-MAR-2003 (revised)

DT 17-AUG-1995 (first entry)

DE Hepatitis C virus (HCV) protease clone C20c DNA.

XX Hepatitis C virus protease; HCV; clone C20c;

XX viral infectivity inhibition; ds.

XX Hepatitis C virus.

XX Key Location/Qualifiers

XX mat\_peptide 1..366

XX /\*tag= a

XX US5371017-A.

XX 06-DEC-1994.

XX 04-APR-1991; 91US-00680296.

XX 04-APR-1990; 90US-00505433.

XX (CHIR ) CHIRON CORP.

XX Houghton M, Choo Q, Kuo G;

XX

DR WPI; 1995-021889/03.

DR P-PSDB; AAR68540.

XX

PT DNA encoding hepatitis C virus protease - used to produce large amts. of  
PT the protease and to develop prods. for inhibition of viral infectivity.

XX

PS Example 3; Fig 2; 69pp; English.

XX

CC AAQ80168 encodes AAR68540 hepatitis C virus (HCV) protease clone C20c,  
CC using recombinant expression systems large amounts of protease can be  
CC produced. The HCV protease can be used in the production of Abs. It can  
CC also be used for assaying agents which inhibit protease activity, to  
CC identify compounds which inhibit viral infectivity. (Updated on 25-MAR-  
CC 2003 to correct Pf field.)

XX

SQ Sequence 368 BP; 70 A; 114 C; 108 G; 76 T; 0 U; 0 Other;

Query Match 2.7%; Score 56; DB 2; Length 368;

Best Local Similarity 100.0%; Pred. No. 1e-17;

Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTGGCGCTCCTCGGACCTTTACCTGTCACGAGGCACGCCGATGTCATTCC 347

Db 220 TGCACCTTGGCGCTCCTCGGACCTTTACCTGTCACGAGGCACGCCGATGTCATTCC 275

RESULT 29

AAT59254

ID AAT59254 standard; DNA; 368 BP.

XX

AC AAT59254;

XX

DT 17-OCT-2003 (revised)

DT 25-MAR-2003 (revised)

DT 03-APR-1997 (first entry)

XX

DE HCV protease clone C20c.

XX

KW HCV; NS3; non-structural domain 3; protease; polyprotein; inhibitor;

XX screen, processing; infection; treatment; probe; hepatitis C virus; ss.

XX

OS Hepatitis C virus; Virus.

XX

PN US5585258-A.

XX

PD 17-DEC-1996.

XX

PF 06-DEC-1994; 94US-00350884.

XX

PR 04-APR-1990; 90US-00505433.

XX

PR 04-APR-1991; 91US-00680296.

XX

PA (CHIR ) CHIRON CORP.

XX

PI Choo Q, Kuo G, Houghton M;

XX

DR WPI; 1997-051175/05.

XX

DR P-PSDB; AAW01690.

XX

PT Compsn. contg. hepatitis C virus NS3 domain protease and related fusion  
PT proteins - useful for screening specific inhibitors, potential antiviral  
PT agents, prepn. of antibodies and for cleaving specific poly:peptide(s).  
XX Example 3; Col 63-64; 68pp; English.  
XX Compsn. comprising the hepatitis C virus (HCV) NS3 domain protease or  
XX its active truncation analogues are claimed. Also new are fusion proteins  
XX comprising the protease (or analogues) and, e.g. human superoxide (SOD)  
XX or ubiquitin. The protease is essential for polyprotein processing, and  
XX thus infectivity, in HCV. The compsns. are used to screen for specific  
XX inhibitors (possibly useful as antiviral agents), to generate specific  
XX antibodies and to cleave specific polypeptides. HCV cDNA clones (AAT59250



```
XX AC ACD44789;
XX AC
XX DT
XX DE
XX DE Hepatitis C virus (HCV) protease clone C20c DNA.
XX DE
XX DE Hepatitis C virus; HCV; protease; protease inhibition; viral infection;
XX KW gene; ds.
XX OS Hepatitis C virus.
XX OS
XX PN US2003027317-A1.
XX PN
XX PD 06-FEB-2003.
XX PF
XX PF 18-JUN-2001; 2001US-00884456.
XX PR
XX PR 04-APR-1990; 90US-00505433.
XX PR 04-APR-1991; 91US-00680296.
XX PR 06-DEC-1994; 94US-00350884.
XX PR 12-MAY-1995; 95US-0040548.
XX PR 06-SEP-1996; 96US-00709177.
XX PR 19-FEB-1999; 99US-00253230.
XX PR
XX (HOUG/) HOUGHTON M.
XX PA (CHOO/) CHOO Q.
XX PA (KUOG/) KUO G.
XX PI
XX PI Houghton M, Choo Q, Kuo G;
XX DR WPI; 2003-492037/58.
XX DR P-PSDB; ABO27013.
XX XX
XX XX New compositions comprising a hepatitis C virus (HCV) protease
XX PT polynucleotide, useful for assaying pharmaceutical agents for controlling
XX PT HCV, and as compounds which inhibit the protease activity and viral
XX PT infectivity.
XX XX
XX PS Example 3; Fig 2; 4lpp; English.
XX XX
XX XX The invention describes a composition comprising a polynucleotide which
XX CC encodes only the hepatitis C virus (HCV) protease or an active HCV
XX CC protease analogue. The protease is useful for assaying pharmaceutical
XX CC agents for controlling HCV, and as compounds which inhibit the protease
XX CC activity sufficiently will also inhibit viral infectivity. This sequence
XX CC encodes hepatitis C virus (HCV) protease
XX SQ Sequence 368 BP; 70 A; 114 C; 108 G; 76 T; 0 U; 0 Other;
Query Match 2.7%; Score 56; DB 8; Length 368;
Best Local Similarity 100.0%; Pred. No. 1e-17;
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 292 TGCACCTTGGCGCTCCTCGGACCTTTACCTGTGCACGAGGACGCCGATGTCATTCC 347
Db 220 TGCACCTTGGCGCTCCTCGGACCTTTACCTGTGCACGAGGACGCCGATGTCATTCC 275
RESULT 33
ADA07860
XX ID ADA07860 standard; cDNA; 368 BP.
XX AC ADA07860;
XX XX
XX DT 06-NOV-2003 (first entry)
XX XX
XX DE Hepatitis C virus cDNA C20c encoding an NS3 protease fragment.
XX KW ss; HCV; virucide; NS3 protease; serine protease; hSOD;
XX KW superoxide dismutase; yeast a-factor; interleukin-2S; ubiquitin;
XX KW beta-galactosidase; beta lactamase; horseradish peroxidase;
XX KW glucose oxidase; urease; HCV infection.
```

```
XX OS Hepatitis C virus.
XX OS
XX PN US2003064499-A1.
XX PN
XX PD 03-APR-2003.
XX PF
XX PF 18-JUN-2001; 2001US-00884455.
XX PR
XX PR 04-APR-1990; 90US-00505433.
XX PR 04-APR-1991; 91US-00680296.
XX PR 06-DEC-1994; 94US-00350884.
XX PR 12-MAY-1995; 95US-0040548.
XX PR 06-SEP-1996; 96US-00709177.
XX PR 18-FEB-1999; 99US-00253675.
XX PR
XX (HOUG/) HOUGHTON M.
XX PA (CHOO/) CHOO Q.
XX PA (KUOG/) KUO G.
XX PI
XX PI Houghton M, Choo Q, Kuo G;
XX DR WPI; 2003-540789/51.
XX DR P-PSDB; ADA07861.
XX XX
XX XX A composition for assaying and designing antiviral agents specific for
XX PT Hepatitis C virus (HCV) comprises a purified proteolytic polypeptide from
XX PT HCV or a polynucleotide which encodes HCV protease.
XX PS
XX PS Example 3; Fig 2; 40pp; English.
XX XX
XX XX The invention relates to a composition comprising a purified proteolytic
XX CC polypeptide derived from Hepatitis C virus (HCV) or a polynucleotide
XX CC which encodes only the HCV protease or an active HCV protease analogue,
XX CC or which encodes a fusion protein comprising HCV protease or HCV protease
XX CC analogue, and a fusion partner. Also included are a fusion protein
XX CC comprising a fusion partner fused to a proteolytic polypeptide derived
XX CC from HCV, a method for assaying compounds for activity against HCV
XX CC (comprising providing an active HCV protease, contacting the protease
XX CC with a compound capable of inhibiting serine protease activity and
XX CC measuring inhibition of the proteolytic activity of the HCV protease) and
XX CC an expression vector for producing HCV protease or HCV protease analogues
XX CC in a host cell (comprising a polynucleotide encoding HCV protease or an
XX CC HCV protease analogue, transcriptional and translational regulatory
XX CC sequences functional in the host cell operably linked to the HCV protease
XX CC -encoding polynucleotide and a selectable marker). The fusion partner is
XX CC selected from hSOD (human superoxide dismutase), yeast a-factor,
XX CC interleukin (IL)-2S, ubiquitin, beta-galactosidase, beta-lactamase,
XX CC horseradish peroxidase, glucose oxidase and urease. The composition is
XX CC useful in assaying and designing antiviral agents specific for HCV. The
XX CC method is used in identifying antiviral agents effective for treating
XX CC HCV. The present sequence is a cDNA encoding an HCV NS3 protease or
XX CC fragment.
XX SQ Sequence 368 BP; 70 A; 114 C; 108 G; 76 T; 0 U; 0 Other;
Query Match 2.7%; Score 56; DB 8; Length 368;
Best Local Similarity 100.0%; Pred. No. 1e-17;
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 292 TGCACCTTGGCGCTCCTCGGACCTTTACCTGTGCACGAGGACGCCGATGTCATTCC 347
Db 220 TGCACCTTGGCGCTCCTCGGACCTTTACCTGTGCACGAGGACGCCGATGTCATTCC 275
RESULT 34
ABX15706
XX ID ABX15706 standard; DNA; 612 BP.
XX AC ABX15706;
XX XX
XX DT 28-MAR-2003 (first entry)
XX XX
```

DE Anti-viral synthetic prototoxophore associated DNA sequence.

XX Hepatitis C; ds; viral prototoxophore; anti-viral; tumour; virus;

KW infection; antitumour; toxophore; human immunodeficiency virus;

KW HIV infection; herpes simplex virus; HSV; rhinovirus; NS3 protease.

XX Unidentified.

XX WO200287500-A2.

PN 07-NOV-2002.

XX 26-APR-2002; 2002WO-US013223.

PF 27-APR-2001; 2001US-0286893P.

PR (NEWB-) NEWBIOTICS INC.

PA Cathers BE, Neuteboom STC, Shepard HW;

XX WPI; 2003-167102/16.

DR Novel synthetic viral prototoxophore for treating viral infections, has

XX toxin moiety incorporated into substrate domain specific for viral

PT enzyme, bound and modified by viral enzyme to get converted into

XX toxophore.

XX Example 1; Page 62; 66pp; English.

XX This invention relates to a novel synthetic viral prototoxophore

CC comprising a toxin moiety operatively incorporated into a substrate

CC domain specific for a viral enzyme. This prototoxophore may be bound and

CC modified by the viral enzyme thus converting it to a toxophore. Also

CC disclosed in the invention is a method for enhancing the anti-viral

CC effect of an antiviral agent, this method comprises contacting a cell,

CC infected with a virus or is susceptible to infection, with a

CC prototoxophore. The invention further comprises an assay to identify anti

CC -viral agents, comprising contacting an infected cell with a candidate

CC agent and comparing the ability of the agent to inhibit the growth or

CC infectivity of the virus in the cell. The prototoxophores of the

CC invention may have virucide or antitumour activity. The prototoxophores

CC of the invention may be useful for reducing or inhibiting viral

CC infectivity, by contacting a cell (e.g. lymphocyte, nerve cell,

CC connective tissue cell, muscle cell or hepatocyte) which is infected with

CC a virus or is susceptible to infection with a virus, with an effective

CC amount of the prototoxophore. The cells are cell lines adapted to long

CC term continuous culture or isolated from a subject. The prototoxophore is

CC also useful for ameliorating the severity of a viral infection in a

CC subject, where the virus is selected from human immunodeficiency virus

CC (HIV), herpes simplex virus (HSV), rhinovirus and hepatitis virus, by

CC administering an effective amount of the prototoxophore to the subject.

CC The prototoxophores of the invention are also useful for treating

CC tumours. The present sequence represents an antiviral prototoxophore

CC associated DNA sequence, this sequence is described as a recombinant

CC NS3/NS4 fusion protein in example 1 of the invention although it is

CC clearly not a protein sequence

XX

XX Sequence 612 BP; 120 A; 171 C; 191 G; 130 T; 0 U; 0 Other;

XX

Query Match 2.7%; Score 56; DB 8; Length 612;

Best Local Similarity 100.0%; Pred. No. 1e-17;

Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTCGGCTCTCTCGGACCTTTACCTGTCACGAGGACCGCGGATGTCATTC 347

349 TGCACCTTCGGCTCTCTCGGACCTTTACCTGTCACGAGGACCGCGGATGTCATTC 404

DB

RESULT 35

AAQ14304

ID AAQ14304 standard; DNA; 1947 BP.

XX

AAQ14304;

XX

Query Match 2.7%; Score 56; DB 8; Length 612;

Best Local Similarity 100.0%; Pred. No. 1e-17;

Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTCGGCTCTCTCGGACCTTTACCTGTCACGAGGACCGCGGATGTCATTC 347

349 TGCACCTTCGGCTCTCTCGGACCTTTACCTGTCACGAGGACCGCGGATGTCATTC 404

DB

RESULT 35

AAQ14304

ID AAQ14304 standard; DNA; 1947 BP.

XX

AAQ14304;

XX

XX 25-MAR-2003 (revised)

DT 14-JAN-1992 (first entry)

XX Vector cf1SODp600 sequence encoding hSOD-HCV protease fusion protein.

DE NANBH; non-A, non-B hepatitis; liver disease;

KW human superoxide dismutase leader sequence; ds.

XX Hepatitis C virus.

OS

XX Key Location/Qualifiers

FT sig\_peptide 1..465

FT /tag= a

FT /label= hSOD leader

FT mat\_peptide 466..1947

FT /tag= b

FT /product= "HCV protease"

XX WO9115575-A.

PN 17-OCT-1991.

XX 04-APR-1990; 90US-00505433.

PF 04-APR-1990; 90US-00505433.

PR (CHIR ) CHIRON CORP.

PA Houghton M, Choo Q, Kuo G;

XX WPI; 1991-325218/44.

DR P-PSDB; AAR14546.

XX New purified protease - derived from hepatitis C virus, for assay, and

PT designing anti-HCV agents.

XX Example 4; Fig 10; 74pp; English.

PS cf1SODp600 was transformed into E.coli D1210 cells and deposited as ATCC

CC 68275. The full-length HCV protease coding sequence was constructed from

CC a number of different clones (see AAQ14297-Q14303). (Updated on 25-MAR-

CC 2003 to correct PA field.)

XX

XX Sequence 1947 BP; 427 A; 539 C; 566 G; 415 T; 0 U; 0 Other;

XX

Query Match 2.7%; Score 56; DB 2; Length 1947;

Best Local Similarity 100.0%; Pred. No. 9.7e-18;

Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTCGGCTCTCTCGGACCTTTACCTGTCACGAGGACCGCGGATGTCATTC 347

1000 TGCACCTTCGGCTCTCTCGGACCTTTACCTGTCACGAGGACCGCGGATGTCATTC 1055

DB

RESULT 36

ABK15344

ID ABK15344 standard; DNA; 2058 BP.

XX

XX ABK15344;

XX

XX 08-MAY-2002 (first entry)

XX

DE Hepatitis C virus NS3/4a conformational epitope gene sequence.

XX Hepatitis C virus; HCV; NS3/4a conformational epitope; seroconversion;

KW immunoassay solid support; multiple epitope fusion antigen; MEPA;

KW non-structural protein; gene; ds.

XX Hepatitis C virus.

XX Key Location/Qualifiers

FT CDS 1..2058

```
FT FT /*tag= a
FT FT /partial
FT FT /product= "HCV NS3/4a conformational epitope"
FT FT /note= "This sequence lacks a stop codon"
XX PN WO200196870-A2.
XX XX 20-DEC-2001.
XX XX 14-JUN-2001; 2001WO-US019156.
XX XX 15-JUN-2000; 2000US-0212082P.
XX XX 02-APR-2001; 2001US-0280811P.
XX XX 02-APR-2001; 2001US-0280867P.
XX PA (CHIR ) CHIRON CORP.
XX PI Chien DY, Arcangel P, Tandeske L, George-Nascimento C, Coit D;
XX PI Medina-Selby A;
XX XX WPI; 2002-090228/12.
XX DR P-PSDB; AAU76377.
XX XX
XX XX Immunoassay solid support, useful for detecting hepatitis C virus
XX PT infection in biological sample, comprises HCV NS3/4a conformational
XX FT epitope and multiple epitope fusion antigen bound to the support.
XX XX
XX PS Disclosure; Fig 3; 92pp; English.
XX CC
XX CC The present invention relates to a new immunoassay solid support
XX CC consisting essentially of at least one hepatitis C virus (HCV) NS3/4a
XX CC conformational epitope and a multiple epitope fusion antigen (MEFA),
XX CC bound to the support. The NS3/4a conformational epitope and/or MEFA
XX CC reacts specifically with anti-HCV antibodies present in a biological
XX CC sample from an HCV-infected individual. The immunoassay of the invention
XX CC is useful for detecting hepatitis C virus infection in a biological
XX CC sample. The method of the invention provides a sensitive, accurate
XX CC diagnostic and prognostic tool to provide adequate patient care and to
XX CC prevent transmission of HCV by blood and by blood products, or by
XX CC personal contact. Use of NS3/4a conformational epitope in combination
XX CC with MEFA, provides a sensitive and reliable method for detecting early
XX CC HCV seroconversion. Use of MEFA has the added advantages of decreasing
XX CC masking problems, improving sensitivity in detecting antibodies by
XX CC allowing a greater number of epitopes on a unit surface area of
XX CC substrate, and improving substrate. Detection accuracy is increased and
XX CC the incidence of false results is reduced because of the identification
XX CC and the use of highly immunogenic HCV antigens which are present during
XX CC the early stages of HCV seroconversion. The present nucleic acid sequence
XX CC encodes the non-structural protein NS3/4a conformational epitope of the
XX CC invention
XX XX
XX SQ Sequence 2058 BP; 419 A; 633 C; 581 G; 425 T; 0 U; 0 Other;
Query Match 2.7%; Score 56; DB 6; Length 2058;
Best Local Similarity 100.0%; Pred. No. 9.7e-18;
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 292 TGCACCTTGGCGCTCCTCGGACCTTTACCTGGTACGAGGACGCGGATGTCATTC 347
Db 292 TGCACCTTGGCGCTCCTCGGACCTTTACCTGGTACGAGGACGCGGATGTCATTC 347
RESULT 37
AAD29795
ID AAD29795 standard; DNA; 2058 BP.
XX AC
XX AC AAD29795;
XX XX
XX DT 17-MAY-2002 (first entry)
XX XX
XX DE HCV-1 NS3/4a mutant conformational antigen encoding DNA.
XX XX
XX KW Hepatitis C virus; NS3/4a antigen; HCV infection; mutant; ds.
```

```
XX XX Hepatitis C virus type 1.
OS Synthetic.
XX XX
XX FH Location/Qualifiers
XX FT 1..686
XX FT /*tag= a
XX FT /product= "HCV-1 NS3/4a conformational antigen"
XX FT /note= "CDS does not include stop codon"
XX FT /partial
XX XX
XX PN WO200196875-A2.
XX XX 20-DEC-2001.
XX XX 14-JUN-2001; 2001WO-US019369.
XX XX 15-JUN-2000; 2000US-0212082P.
XX XX 02-APR-2001; 2001US-0280811P.
XX XX 02-APR-2001; 2001US-0280867P.
XX XX (CHIR ) CHIRON CORP.
XX PA
XX PI Chien DY, Arcangel P, Tandeske L, George-Nascimento C, Coit D;
XX PI Medina-Selby A;
XX XX WPI; 2002-179522/23.
XX DR P-PSDB; AAE18689.
XX XX
XX PT Immunoassay solid support useful for detecting hepatitis C virus
XX PT infection in a biological sample, comprises at least one of HCV anti-core
XX FT antibody and HCV NS3/4a epitope, bound to the support.
XX XX
XX PS Example 2; Fig 4; 87pp; English.
XX CC
XX CC The present invention relates to hepatitis C virus (HCV) core antigen and
XX CC NS (nonstructural) 3/4a antibody combination assay that can detect both
XX CC HCV antigens and antibodies present in a sample using a single solid
XX CC matrix as well as immunoassay solid supports for use in the assay. The
XX CC solid support is useful for detecting HCV infection in a biological
XX CC sample. The present sequence is a DNA encoding HCV-1 NS3/4a mutant
XX CC conformational antigen. This sequence is used in the exemplification of
XX CC the invention
XX XX
XX SQ Sequence 2058 BP; 419 A; 634 C; 580 G; 425 T; 0 U; 0 Other;
Query Match 2.7%; Score 56; DB 6; Length 2058;
Best Local Similarity 100.0%; Pred. No. 9.7e-18;
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 292 TGCACCTTGGCGCTCCTCGGACCTTTACCTGGTACGAGGACGCGGATGTCATTC 347
Db 292 TGCACCTTGGCGCTCCTCGGACCTTTACCTGGTACGAGGACGCGGATGTCATTC 347
RESULT 38
ABX14410
ID ABX14410 standard; DNA; 2058 BP.
XX AC
XX AC ABX14410;
XX XX
XX DT 06-MAR-2003 (first entry)
XX XX
XX DE DNA encoding HCV-1 NS3/4a conformational antigen.
XX XX
XX KW Immunoassay solid support; Hepatitis C Virus type-1; HCV-1;
XX KW NS3/4a conformational epitope; multiple epitope fusion antigen; MEFA;
XX KW anti-HCV antibody; NS3/4a conformational antigen; HCV infection; mutant;
XX KW gene; ds.
XX XX
XX OS Hepatitis C virus type 1.
XX OS Synthetic.
XX XX
```

KW non-A, non-B hepatitis; NANB; conformational epitope; mutant; ds; gene.  
 XX Synthetic.  
 OS Hepatitis C virus.  
 XX Key Location/Qualifiers  
 FT 1..2058  
 FT /\*tag= a  
 FT /partial  
 FT /product= "NS3/4a conformational antigen"  
 FT /note= "This sequence lacks a stop codon"

XX US2002146685-A1.

XX 10-OCT-2002.

XX 14-JUN-2001; 2001US-00881654.

XX 15-JUN-2000; 2000US-0212082P.

XX 02-APR-2001; 2001US-0280811P.

XX 02-APR-2001; 2001US-0280867P.

XX (CHIE//) CHIEN D Y.

XX (AFCA//) ARCANGEL P.

XX (TAND//) TANDESKE L.

XX (GEOR//) GEORGE-NASCIMENTO C.

XX (COIT//) COIT D.

XX (MEDI//) MEDINA-SELBY A.

XX Chien DY, Arcangel P, Tandeske L, George-Nascimento C, Coit D;

XX Medina-Selby A;

XX WPI; 2003-147573/14.

XX P-PSDB; ABG72261.

XX Immunoassay solid support for detecting Hepatitis C Virus infection in biological samples, comprises Hepatitis C Virus conformational epitope and multiple epitope fusion antigen.

XX Disclosure; Fig 3A-3D; 45pp; English.

XX The present invention relates to immunoassays comprising Hepatitis C Virus (HCV) NS3/4a conformational epitope and multiple epitope fusion antigen (MEFA), bound to a solid support. The NS3/4a epitope and/or the multiple epitope fusion antigen react with anti-HCV antibodies present in a biological sample from an HCV-infected individual. The immunoassays and methods of the invention are useful for detecting HCV infection in a biological sample. The inventive immunoassay solid support provides a sensitive and reliable method for detecting early HCV seroconversion. The assays can detect HCV infection caused by any six known genotypes of HCV. The use of the multiple epitope fusion proteins decreases masking problems, improves sensitivity in detecting antibodies by allowing a greater number of epitopes on a unit area of substrate, and improves selectivity. The present sequence encodes HCV type 1 (HCV-1) NS3/4a conformational antigen, a mutant of the HCV-1 NS3/4a polypeptide

XX Sequence 2058 BP; 419 A; 633 C; 581 G; 425 T; 0 U; 0 Other;

XX Query Match 2.7%; Score 56; DB 7; Length 2058;

XX Best Local Similarity 100.0%; Pred.No. 9.7e-18;

XX Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTGGCGCTCCTCGGACCTTTACCTGGTCACGAGGACCGCGATGTCATCC 347

DB 292 TGCACCTTGGCGCTCCTCGGACCTTTACCTGGTCACGAGGACCGCGATGTCATCC 347

RESULT 39

ADCO6768

ID ADC06768 standard; DNA; 2058 BP.

XX AC ADC06768;

XX 18-DEC-2003 (first entry)

XX HCV mutant conformational NS3/4a epitope DNA.

XX immunoassay solid support; HCV; NS3/4a; non-structural;

KW non-A, non-B hepatitis; NANB; conformational epitope; mutant; ds; gene.  
 XX Synthetic.  
 OS Hepatitis C virus.  
 XX Key Location/Qualifiers  
 FT 1..2058  
 FT /\*tag= a  
 FT /partial  
 FT /product= "HCV mutant conformational NS3/4a epitope  
 FT protein T403P/S404I"  
 FT /note= "No stop codon"

XX US2002192639-A1.

XX 19-DEC-2002.

XX 14-JUN-2001; 2001US-00881239.

XX 15-JUN-2000; 2000US-0212082P.

XX 02-APR-2001; 2001US-0280811P.

XX 02-APR-2001; 2001US-0280867P.

XX (CHIE//) CHIEN D Y.

XX (ARCA//) ARCANGEL P.

XX (TAND//) TANDESKE L.

XX (GEOR//) GEORGE-NASCIMENTO C.

XX (COIT//) COIT D.

XX (MEDI//) MEDINA-SELBY A.

XX Chien DY, Arcangel P, Tandeske L, George-Nascimento C, Coit D;

XX Medina-Selby A;

XX WPI; 2003-644509/61.

XX P-PSDB; ABC06767.

XX Immunoassay solid support for detecting hepatitis C virus infection in biological samples, comprises a hepatitis C virus anti-core antibody and an isolated hepatitis C virus NS3/4a epitope bound HCV anti-core antibody.

XX Example 2; Fig 4; 40pp; English.

XX The invention relates to a novel immunoassay solid support comprising at least one hepatitis C virus (HCV) anti-core antibody and at least one isolated HCV NS3/4a (non-structural protein 3/4a) epitope bound thereto. The system of the invention may be useful for detecting HCV infection in a biological sample and for treating or detecting non-A, non-B hepatitis (NANB hepatitis). The current sequence is that of the HCV mutant conformational NS3/4a epitope DNA of the invention.

XX Sequence 2058 BP; 419 A; 634 C; 580 G; 425 T; 0 U; 0 Other;

XX Query Match 2.7%; Score 56; DB 9; Length 2058;

XX Best Local Similarity 100.0%; Pred.No. 9.7e-18;

XX Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTGGCGCTCCTCGGACCTTTACCTGGTCACGAGGACCGCGATGTCATCC 347

DB 292 TGCACCTTGGCGCTCCTCGGACCTTTACCTGGTCACGAGGACCGCGATGTCATCC 347

RESULT 40

AAQ14296

ID AAQ14296 standard; cDNA; 2064 BP.

XX AC AAQ14296;

XX 25-MAR-2003 (revised)

XX 14-JAN-1992 (first entry)

XX Hepatitis C Virus protease.

KW NANBH; non-A, non-B hepatitis; fusion protein; ds.  
XX Hepatitis C virus.  
XX  
XX  
XX Key Location/Qualifiers  
FT CDS 7..2064  
FT /\*tag= a  
FT /product= "HCV protease"  
XX  
XX WO9115575-A.  
XX  
XX 17-OCT-1991.  
XX  
XX 04-APR-1990; 90US-00505433.  
XX  
XX 04-APR-1990; 90US-00505433.  
XX  
XX (CHIR ) CHIRON CORP.  
XX  
XX Houghton M, Choo Q, Kuo G;  
XX  
XX WPI; 1991-325218/44.  
XX P-PSDB; AAR14538.  
XX  
XX New purified protease - derived from hepatitis C virus, for assay, and  
XX designing anti-HCV agents.  
XX  
XX Claim 5; Fig 1; 74pp; English.  
XX  
XX HCV protease is encoded in the NS3 domain of the HCV genome. The sequence  
XX varies from strain to strain. The recombinant protease is preferably  
XX produced as a fusion protein in which the fusion partner is human  
XX superoxide dismutase, yeast alpha-factor, IL-2 signal, ubiquitin, beta-  
XX galactosidase, beta-lactamase, horseradish peroxidase, glucose oxidase or  
XX urease. See also AAQ14297-Q14311. (Updated on 25-MAR-2003 to correct PA  
XX field.)  
XX  
XX Sequence 2064 BP; 413 A; 640 C; 584 G; 427 T; 0 U; 0 Other;  
XX  
XX Query Match 2.7%; Score 56; DB 2; Length 2064;  
XX Best Local Similarity 100.0%; Pred. No. 9.7e-18;  
XX Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
XX  
XX QY 292 TGCACCTTGGCGCTCCTCGGACCTTTACCTGTCACGAGGCACGCCGATGTCATCC 347  
XX 541 TGCACCTTGGCGCTCCTCGGACCTTTACCTGTCACGAGGCACGCCGATGTCATCC 596  
XX  
XX  
XX RESULT 41  
XX AAQ80167  
XX ID AAQ80167 standard; DNA; 2064 BP.  
XX  
XX AC AAQ80167;  
XX  
XX 25-MAR-2003 (revised)  
XX 16-AUG-1995 (first entry)  
XX  
XX Hepatitis C virus (HCV) protease DNA.  
XX  
XX Hepatitis C virus protease; HCV; viral infectivity inhibition; ds.  
XX  
XX Hepatitis C virus.  
XX  
XX Key Location/Qualifiers  
XX mat\_peptide 7..2064  
XX /\*tag= a  
XX  
XX US5371017-A.  
XX  
XX 06-DEC-1994.  
XX  
XX 04-APR-1991; 91US-00680296.  
XX  
XX

PR 04-APR-1990; 90US-00505433.  
XX  
XX (CHIR ) CHIRON CORP.  
XX  
XX Houghton M, Choo Q, Kuo G;  
XX  
XX WPI; 1995-021889/03.  
XX P-PSDB; AAR66267.  
XX  
XX DNA encoding hepatitis C virus protease - used to produce large ants. of  
XX the protease and to develop prods. for inhibition of viral infectivity.  
XX  
XX Claim 2; Fig 1; 69pp; English.  
XX  
XX AAQ80167 encodes AAR66267 hepatitis C virus (HCV) protease, using  
XX recombinant expression systems large amounts of protease can be produced.  
XX The HCV protease can be used in the production of Abs. It can also be  
XX used for assaying agents which inhibit protease activity, to identify  
XX compounds which inhibit viral infectivity. (Updated on 25-MAR-2003 to  
XX correct PF field.)  
XX  
XX Sequence 2064 BP; 413 A; 641 C; 583 G; 427 T; 0 U; 0 Other;  
XX  
XX Query Match 2.7%; Score 56; DB 2; Length 2064;  
XX Best Local Similarity 100.0%; Pred. No. 9.7e-18;  
XX Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
XX  
XX QY 292 TGCACCTTGGCGCTCCTCGGACCTTTACCTGTCACGAGGCACGCCGATGTCATCC 347  
XX 541 TGCACCTTGGCGCTCCTCGGACCTTTACCTGTCACGAGGCACGCCGATGTCATCC 596  
XX  
XX  
XX RESULT 42  
XX AAT59260  
XX ID AAT59260 standard; DNA; 2064 BP.  
XX  
XX AC AAT59260;  
XX  
XX 17-OCT-2003 (revised)  
XX 25-MAR-2003 (revised)  
XX 03-APR-1997 (first entry)  
XX  
XX DNA encoding Hepatitis C virus NS3 domain protease.  
XX  
XX HCV; NS3; non-structural domain 3; protease; polypeptide; inhibitor;  
XX screen; processing; infection; treatment; ss.  
XX  
XX Hepatitis C virus; Virus.  
XX  
XX Key Location/Qualifiers  
XX CDS 7..2064  
XX /\*tag= a  
XX /note= "NS3 domain polypeptide"  
XX  
XX US5585258-A.  
XX  
XX 17-DEC-1996.  
XX  
XX 06-DEC-1994; 94US-00350884.  
XX  
XX 04-APR-1990; 90US-00505433.  
XX  
XX 04-APR-1991; 91US-00680296.  
XX  
XX (CHIR ) CHIRON CORP.  
XX  
XX Choo Q, Kuo G, Houghton M;  
XX  
XX WPI; 1997-051175/05.  
XX P-PSDB; AAW01693.  
XX  
XX Compn. contg. hepatitis C virus NS3 domain protease and related fusion  
XX proteins - useful for screening specific inhibitors, potential antiviral  
XX agents, prepn. of antibodies and for cleaving specific poly:peptide(s).  
XX  
XX

XX PS Example 1; Col 53-60; 68pp; English.

XX CC This sequence encodes part of the hepatitis C virus (HCV) polyprotein

CC CC which contains the NS3 domain protease. Compsns. comprising the HCV

CC CC protease or its active truncation analogues are claimed. Also new are

CC CC fusion proteins comprising the protease (or analogues) and, e.g. human

CC CC superoxide or ubiquitin. The protease is essential for polyprotein

CC CC processing, and thus infectivity, in HCV. The compsns. are used to screen

CC CC for specific inhibitors (possibly useful as antiviral agents), to

CC CC generate specific antibodies and to cleave specific polypeptides.

CC CC (Updated on 25-MAR-2003 to correct PF field.) (Updated on 17-OCT-2003 to

CC CC standardise OS field)

XX SQ Sequence 2064 BP; 413 A; 641 C; 583 G; 427 T; 0 U; 0 Other;

Query Match 2.7%; Score 56; DB 2; Length 2064;  
Best Local Similarity 100.0%; Pred. No. 9.7e-18;  
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTGGCGCTCCTCGACCTTTACCTGCTCAGGACGCGCGGATGTCATTC 347  
|||||

Db 541 TGCACCTTGGCGCTCCTCGACCTTTACCTGCTCAGGACGCGCGGATGTCATTC 596  
|||||

RESULT 43

AAV04985

ID AAV04985 standard; DNA; 2064 BP.

XX AC AAV04985;

XX DT 27-AUG-2003 (revised)

XX DT 07-MAY-1998 (first entry)

XX DE Nucleotide sequence encoding Hepatitis C virus (HCV) protease.

XX KW Protease; HCV; NS3 domain; human superoxide dismutase; fusion protein;

XX KW ubiquitin; assay; activity; anti-HCV; ss.

XX OS Hepatitis C virus.

Key Location/Qualifiers

CDS 7..2064

FT /\*tag= a

FT /note= "no ATG start or STOP codons given"

XX US5712145-A.

XX PD 27-JAN-1998.

XX PF 06-SEP-1996; 96US-00709173.

XX PR 04-APR-1990; 90US-00505433.

XX PR 04-APR-1991; 91US-00680296.

XX PR 06-DEC-1994; 94US-00350884.

XX PR 12-MAY-1995; 95US-00440548.

XX PA (CHIR ) CHIRON CORP.

XX PI Choo Q, Kuo G, Houghton M;

XX DR WPI; 1998-119986/11.

XX DR P-PSDB; AAW46389.

XX PT Recombinant hepatitis C virus protease - useful in screening drugs for

XX PT activity against hepatitis C virus.

XX PS Disclosure; Fig 1A-F; 68pp; English.

XX CC The present sequence encodes a Hepatitis C virus (HCV) protease. The

CC CC protein is encoded by sequences within the NS3 domain. The protease is

CC CC believed to cleave itself from the genomic polyprotein. In the absence of

CC CC protease activity, the HCV polyprotein should remain in its unprocessed

CC form, and thus render the virus non-infectious. Inhibitors of protease

CC activity should therefore also inhibit viral infectivity. The protease is

CC used for assaying compounds for activity against HCV. (Updated on 27-AUG-

CC CC 2003 to correct OS field.)

XX SQ Sequence 2064 BP; 413 A; 641 C; 583 G; 427 T; 0 U; 0 Other;

Query Match 2.7%; Score 56; DB 2; Length 2064;  
Best Local Similarity 100.0%; Pred. No. 9.7e-18;  
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTGGCGCTCCTCGACCTTTACCTGCTCAGGACGCGCGGATGTCATTC 347  
|||||

Db 541 TGCACCTTGGCGCTCCTCGACCTTTACCTGCTCAGGACGCGCGGATGTCATTC 596  
|||||

RESULT 44

AAAX26390

ID AAX26390 standard; DNA; 2064 BP.

XX AC AAX26390;

XX DT 26-MAY-1999 (first entry)

XX DE DNA encoding the HCV protease of the invention.

XX KW HCV NS3 protease; truncation analog; HCV control; protease activity;

XX KW viral infectivity; inactive non-cleaving protease; ss.

XX OS Hepatitis C virus.

Key Location/Qualifiers

CDS 7..2064

FT /\*tag= a

FT /note= "no termination codon"

XX US5885799-A.

XX PD 23-MAR-1999.

XX PF 06-SEP-1996; 96US-00709177.

XX PR 04-APR-1990; 90US-00505433.

XX PR 04-APR-1991; 91US-00680296.

XX PR 06-DEC-1994; 94US-00350884.

XX PR 12-MAY-1995; 95US-00440548.

XX PA (CHIR ) CHIRON CORP.

XX PI Choo Q, Kuo G, Houghton M;

XX DR WPI; 1999-228536/19.

XX DR P-PSDB; AAW97601.

XX PT Preparation of new Hepatitis C Virus NS3 protease - useful for screening

XX PT for compounds which inhibit HCV infectivity.

XX PS Example 4; Fig 1; 71pp; English.

XX CC The specification describes a method for making a purified Hepatitis C

CC CC virus (HCV) NS3 protease or active truncation analog. If the HCV protease

CC CC N-terminal cleavage signal is excluded (so that self-cleavage is

CC CC prevented), the HCV protease remains in its unprocessed form, and renders

CC CC the virus noninfectious. The protease is therefore useful for assaying

CC CC pharmaceutical agents for control of HCV, as compounds which inhibit

CC CC protease activity sufficiently will also inhibit viral infectivity. An

CC CC inactive non-cleaving protease can be used to screen for inhibitors.

CC CC Recombinant expression systems can be utilised to prepare recombinant HCV

CC CC which can be used to produce monoclonal antibodies. The present sequence

CC CC appears in the specification

XX SQ Sequence 2064 BP; 413 A; 641 C; 583 G; 427 T; 0 U; 0 Other;



Query Match 2.7%; Score 56; DB 2; Length 2064;  
 Best Local Similarity 100.0%; Pred. No. 9.7e-18;  
 Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 292 TGCACCTTGGCGCTCTCGGACCTTTACCTGGTCAAGGACGCGCGATGTCATTC 347  
 Db 541 TGCACCTTGGCGCTCTCGGACCTTTACCTGGTCAAGGACGCGCGATGTCATTC 596

RESULT 45  
 ACD44788  
 ID ACD44788 standard; DNA; 2064 BP.  
 XX ACD44788;  
 XX  
 DT 09-SEP-2003 (first entry)  
 XX  
 DE Hepatitis C virus (HCV) protease DNA.  
 XX  
 KW Hepatitis C virus; HCV; protease; protease inhibition; viral infection;  
 KW gene; ds.  
 XX  
 OS Hepatitis C virus.  
 XX  
 PN US2003027317-A1.  
 XX  
 PD 06-FEB-2003.  
 XX  
 PF 18-JUN-2001; 2001US-00884456.  
 XX  
 PR 04-APR-1990; 90US-00505433.  
 PR 04-APR-1991; 91US-00680296.  
 PR 06-DEC-1994; 94US-00350884.  
 PR 12-MAY-1995; 95US-00440548.  
 PR 06-SEP-1996; 96US-00709177.  
 PR 19-FEB-1999; 99US-00253230.  
 XX  
 PA (HOUG/) HOUGHTON M.  
 PA (CHOO/) CHOO Q.  
 PA (KUOG/) KUO G.  
 XX  
 PI Houghton M, Choo Q, Kuo G;  
 XX  
 DR WPI; 2003-492037/58.  
 DR P-PSDB; ABO27012.  
 XX  
 PT New compositions comprising a hepatitis C virus (HCV) protease  
 PT polynucleotide, useful for assaying pharmaceutical agents for controlling  
 PT HCV, and as compounds which inhibit the protease activity and viral  
 PT infectivity.  
 XX  
 PS Example 2; Fig 1; 41pp; English.  
 XX  
 CC The invention describes a composition comprising a polynucleotide which  
 CC encodes only the hepatitis C virus (HCV) protease or an active HCV  
 CC protease analogue. The protease is useful for assaying pharmaceutical  
 CC agents for controlling HCV, and as compounds which inhibit the protease  
 CC activity sufficiently will also inhibit viral infectivity. This sequence  
 CC encodes hepatitis C virus (HCV) protease  
 XX  
 SQ Sequence 2064 BP; 413 A; 641 C; 583 G; 427 T; 0 U; 0 Other;  
 Query Match 2.7%; Score 56; DB 8; Length 2064;  
 Best Local Similarity 100.0%; Pred. No. 9.7e-18;  
 Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 292 TGCACCTTGGCGCTCTCGGACCTTTACCTGGTCAAGGACGCGCGATGTCATTC 347  
 Db 541 TGCACCTTGGCGCTCTCGGACCTTTACCTGGTCAAGGACGCGCGATGTCATTC 596

RESULT 46  
 ADA07858

ID ADA07858 standard; cDNA; 2064 BP.  
 XX  
 AC ADA07858;  
 XX  
 DT 06-NOV-2003 (first entry)  
 XX  
 DE Hepatitis C virus cDNA encoding the NS3 protease.  
 XX  
 KW ss; gene; HCV; virucide; NS3 protease; serine protease; hSOD;  
 KW superoxide dismutase; yeast a-factor; interleukin-2S; ubiquitin;  
 KW beta-galactosidase; beta-lactamase; horseradish peroxidase;  
 KW glucose oxidase; urease; HCV infection.  
 XX  
 OS Hepatitis C virus.  
 XX  
 PN US2003064499-A1.  
 XX  
 PD 03-APR-2003.  
 XX  
 PF 18-JUN-2001; 2001US-00884455.  
 XX  
 PR 04-APR-1990; 90US-00505433.  
 PR 04-APR-1991; 91US-00680296.  
 PR 06-DEC-1994; 94US-00350884.  
 PR 12-MAY-1995; 95US-00440548.  
 PR 06-SEP-1996; 96US-00709177.  
 PR 18-FEB-1999; 99US-00253675.  
 XX  
 PA (HOUG/) HOUGHTON M.  
 PA (CHOO/) CHOO Q.  
 PA (KUOG/) KUO G.  
 XX  
 PI Houghton M, Choo Q, Kuo G;  
 XX  
 DR WPI; 2003-540789/51.  
 DR P-PSDB; ADA07859.  
 XX  
 PT A composition for assaying and designing antiviral agents specific for  
 PT Hepatitis C virus (HCV) comprises a purified proteolytic polypeptide from  
 PT HCV or a polynucleotide which encodes HCV protease.  
 XX  
 PS Example 2; Fig 1; 40pp; English.  
 XX  
 CC The invention relates to a composition comprising a purified proteolytic  
 CC polypeptide derived from Hepatitis C virus (HCV) or a polynucleotide  
 CC which encodes only the HCV protease or an active HCV protease analogue,  
 CC or which encodes a fusion protein comprising HCV protease or HCV protease  
 CC analogue, and a fusion partner. Also included are a fusion protein  
 CC comprising a fusion partner fused to a proteolytic polypeptide derived  
 CC from HCV, a method for assaying compounds for activity against HCV  
 CC with a compound capable of inhibiting serine protease activity and  
 CC measuring inhibition of the proteolytic activity of the HCV protease) and  
 CC an expression vector for producing HCV protease or HCV protease analogues  
 CC in a host cell (comprising a polynucleotide encoding HCV protease or an  
 CC HCV protease analogue, transcriptional and translational regulatory  
 CC sequences functional in the host cell operably linked to the HCV protease  
 CC -encoding polynucleotide and a selectable marker). The fusion partner is  
 CC selected from hSOD (human superoxide dismutase), yeast a-factor,  
 CC interleukin (IL)-2S, ubiquitin, beta-galactosidase, beta-lactamase,  
 CC horseradish peroxidase, glucose oxidase and urease. The composition is  
 CC useful in assaying and designing antiviral agents specific for HCV. The  
 CC method is used in identifying antiviral agents effective for treating  
 CC HCV. The present sequence is the full length cDNA encoding the HCV NS3  
 CC protease.  
 XX  
 SQ Sequence 2064 BP; 413 A; 641 C; 583 G; 427 T; 0 U; 0 Other;  
 Query Match 2.7%; Score 56; DB 8; Length 2064;  
 Best Local Similarity 100.0%; Pred. No. 9.7e-18;  
 Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 292 TGCACCTTGGCGCTCTCGGACCTTTACCTGGTCAAGGACGCGCGATGTCATTC 347

Db 541 TGCACCTTGGCTCTCGGACCTTTACCTGTCACGAGGACCGCGATGTCATTCC 596  
|||||

RESULT 47  
ABX15705  
ID ABX15705 standard; cDNA; 2073 BP.  
XX AC ABX15705;  
XX DT 28-MAR-2003 (first entry)  
XX DE cDNA sequence encoding hepatitis C virus NS3 and NS4A protease genes.  
XX KW Hepatitis C; ss; viral prototoxophore; anti-viral; tumour; NS4A; virus;  
XX KW infection; antitumour; toxophore; human immunodeficiency virus;  
XX KW HIV infection; herpes simplex virus; HSV; rhinovirus; NS3 protease.  
XX OS Hepatitis C virus.  
XX OS Synthetic.  
XX PN WO200287500-A2.  
XX PD 07-NOV-2002.  
XX PF 26-APR-2002; 2002WO-US013223.  
XX PR 27-APR-2001; 2001US-0286893P.  
XX PA (NEWB-) NEWBIOTICS INC.  
XX PI Cathers BE, Neuteboom STC, Shepard HM;  
XX WPI; 2003-167102/16.  
XX DR Novel synthetic viral prototoxophore for treating viral infections, has  
XX PT toxin moiety incorporated into substrate domain specific for viral  
XX PT enzyme, bound and modified by viral enzyme to get converted into  
XX PT toxophore.  
XX PS Example 1; Page 62; 66pp; English.

This invention relates to a novel synthetic viral prototoxophore comprising a toxin moiety operatively incorporated into a substrate domain specific for a viral enzyme. This prototoxophore may be bound and modified by the viral enzyme thus converting it to a toxophore. Also disclosed in the invention is a method for enhancing the anti-viral effect of an antiviral agent, this method comprises contacting a cell, infected with a virus or is susceptible to infection, with a prototoxophore. The invention further comprises an assay to identify anti-viral agents, comprising contacting an infected cell with a candidate agent and comparing the ability of the agent to inhibit the growth or infectivity of the virus in the cell. The prototoxophores of the invention may have virucide or antitumour activity. The prototoxophores of the invention may be useful for reducing or inhibiting viral infectivity, by contacting a cell (e.g. lymphocyte, nerve cell, connective tissue cell, muscle cell or hepatocyte) which is infected with a virus or is susceptible to infection with a virus, with an effective amount of the prototoxophore. The cells are cell lines adapted to long term continuous culture or isolated from a subject. The prototoxophore is also useful for ameliorating the severity of a viral infection in a subject, where the virus is selected from human immunodeficiency virus (HIV), herpes simplex virus (HSV), rhinovirus and hepatitis virus, by administering an effective amount of the prototoxophore to the subject. The prototoxophores of the invention are also useful for treating tumours. The present sequence represents a cDNA sequence encoding the Hepatitis C virus NS3 and NS4A protease genes for use in an assay to measure the conversion of a prototoxophore of the invention to an activated toxophore in cells transfected with this gene

XX SQ Sequence 2073 BP; 423 A; 635 C; 581 G; 434 T; 0 U; 0 Other;  
Query Match 2.7%; Score 56; DB 8; Length 2073;  
XX

Best Local Similarity 100.0%; Pred. No. 9.7e-18;  
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 292 TGCACCTTGGCTCTCGGACCTTTACCTGTCACGAGGACCGCGATGTCATTCC 347  
|||||  
Db 298 TGCACCTTGGCTCTCGGACCTTTACCTGTCACGAGGACCGCGATGTCATTCC 353  
|||||

RESULT 48  
AAQ14358  
ID AAQ14358 standard; DNA; 2523 BP.  
XX AC AAQ14358;  
XX DT 16-JAN-1992 (first entry)  
XX DE HCV protease gene:hsod leader fusion in cflSODp600.  
XX KW Hepatitis C virus; HCV; human superoxide dismutase; SOD; ss.  
XX OS Hepatitis C virus.  
XX FH Key  
XX FT Signal\_peptide 1..465  
XX FT /tag= a  
XX FT /note= "hsod leader sequence"  
XX FT 466..2523  
XX FT /tag= b  
XX FT /product= "AAS 946-1630 of HCV"  
XX PN WO9115596-A.  
XX PD 17-OCT-1991.  
XX PF 04-APR-1990; 90US-00505434.  
XX PR 04-APR-1990; 90US-00505434.  
XX PA (PROT-) PROTOS INC.  
XX PI Rosenberg S;  
XX DR ~~WPI; 1991-325236/44.~~  
XX DR P-PSDB; AAR14349.  
XX PT Method for assaying pharmaceutical cpds. - for determining anti-Hepatitis  
XX C Virus activity, using binding affinity.  
XX PS Example 4; Fig 10; 68pp; English.

The vector cflSODp600 contains a full-length HCV protease coding sequence fused to a functional hsod leader. The truncated protease analogue expressed by the vector is proteolytically inactive and can be used to assay a wide range of pharmaceutical agents for controlling HCV. Those agents which inhibit the protease activity sufficiently will also inhibit viral infectivity. See also AAQ14359-Q14365  
XX SQ Sequence 2523 BP; 546 A; 722 C; 718 G; 537 T; 0 U; 0 Other;  
Query Match 2.7%; Score 56; DB 2; Length 2523;  
Best Local Similarity 100.0%; Pred. No. 9.7e-18;  
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 292 TGCACCTTGGCTCTCGGACCTTTACCTGTCACGAGGACCGCGATGTCATTCC 347  
|||||  
Db 1000 TGCACCTTGGCTCTCGGACCTTTACCTGTCACGAGGACCGCGATGTCATTCC 1055  
|||||

RESULT 49  
AAQ80175  
ID AAQ80175 standard; DNA; 2523 BP.  
XX AC AAQ80175;  
Query Match 2.7%; Score 56; DB 8; Length 2073;  
XX

same as 5-885, 789A

XX 25-MAR-2003 (revised)  
DT 17-AUG-1995 (first entry)  
XX HCV protease/hSOD fusion protein expression vector cflSODp600.  
XX Hepatitis C virus protease/hSOD fusion protein; HCV;  
KW expression vector cflSODp600; viral infectivity inhibition; ds.  
XX Hepatitis C virus.  
OS  
XX Key Location/Qualifiers  
FH CDS 1. .2523  
FT FT /tag= a  
FT FT 1. .465  
FT FT /tag= b  
FT FT /label= hSOD leader  
XX  
XX US5371017-A.  
XX 06-DEC-1994.  
XX 04-APR-1991; 91US-00680296.  
XX 04-APR-1990; 90US-00505433.  
XX (CHIR ) CHIRON CORP.  
XX Houghton M, Choo Q, Kuo G;  
XX WPI; 1995-021889/03.  
DR P-PSDB; AAR68547.  
XX DNA encoding hepatitis C virus protease - used to produce large amts. of  
PT the protease and to develop prods. for inhibition of viral infectivity.  
XX Claim 10; Fig 10; 69pp; English.  
XX AAQ80175 (which encodes AAR68547) describes the sequence of the hepatitis  
CC C virus (HCV) protease/hSOD fusion protein E. coli expression vector,  
CC cflSODp600. Other claimed HCV protease fusion partners are yeast alpha-  
CC factor, IL-28, ubiquitin, beta-galactosidase, beta-lactamase, horseradish  
CC peroxidase, glucose oxidase and urease. The HCV protease fusion proteins  
CC can be used in the production of Abs. They can also be used for assaying  
CC agents which inhibit protease activity, to identify compounds which  
CC inhibit viral infectivity. (Updated on 25-MAR-2003 to correct PF field.)  
XX  
SQ Sequence 2523 BP; 544 A; 724 C; 718 G; 537 T; 0 U; 0 Other;  
Query Match 2.7%; Score 56; DB 2; Length 2523;  
Best Local Similarity 100.0%; Pred. No. 9.7e-18;  
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 292 TGCACCTTGGCGCTCTCTCGACCTTTACCTGGTCAAGGACGCGCGATGTCATTCC 347  
DB 1000 TGCACCTTGGCGCTCTCTCGACCTTTACCTGGTCAAGGACGCGCGATGTCATTCC 1055  
RESULT 50  
AAT59261  
ID AAT59261 standard; DNA; 2523 BP.  
XX  
AC AAT59261;  
XX  
DT 17-OCT-2003 (revised)  
DT 25-MAR-2003 (revised)  
DT 03-APR-1997 (first entry)  
XX  
DE cflSODp600 encoding hSOD-HCV fusion protein.  
XX HCV; NS3; non-structural domain 3; protease; polyprotein; inhibitor;  
KW screen; processing; infection; treatment; probe; hepatitis C virus; ss.  
XX

OS Hepatitis C virus; Virus.  
OS Homo sapiens.  
OS Chimeric.  
XX  
PN US5585258-A.  
XX 17-DEC-1996.  
XX 06-DEC-1994; 94US-00350884.  
XX 04-APR-1990; 90US-00505433.  
XX 04-APR-1991; 91US-00680296.  
XX (CHIR ) CHIRON CORP.  
XX Choo Q, Kuo G, Houghton M;  
XX WPI; 1997-051175/05.  
XX P-PSDB; AAW01701.  
XX Compsn. contg. hepatitis C virus NS3 domain protease and related fusion  
PT proteins - useful for screening specific inhibitors, potential antiviral  
PT agents, prepn. of antibodies and for cleaving specific poly(peptide)s.  
XX Example 4; Col 77-84; 68pp; English.  
XX Compsns. comprising the hepatitis C virus (HCV) NS3 domain protease or  
CC its active truncation analogues are claimed. Also new are fusion proteins  
CC comprising the protease (or analogues) and, e.g. human superoxide (SOD)  
CC or ubiquitin. The protease is essential for polypeptide processing, and  
CC thus infectivity, in HCV. The compsns. are used to screen for specific  
CC inhibitors (possibly useful as antiviral agents), to generate specific  
CC antibodies and to cleave specific polypeptides. HCV cDNA clones (AAT59250  
CC - 56 encoding AAW01686-92 resp.) were isolated from HCV genomic library  
CC using probes AAT59244-49. The clones were used in the preparation of full  
CC length SOD-protease fusion proteins. The present sequence is that of  
CC vector cflSODp600 which contains a full-length HCV protease coding  
CC sequence fused to a functional hSOD leader. The resulting vector encodes  
CC amino acids 1-151 of hSOD, and amino acids 946-1630 of HCV (corresponding  
CC to 1-686 of AAW01693). (Updated on 25-MAR-2003 to correct PF field.)  
XX (Updated on 17-OCT-2003 to standardise OS field)  
SQ Sequence 2523 BP; 545 A; 722 C; 719 G; 537 T; 0 U; 0 Other;  
Query Match 2.7%; Score 56; DB 2; Length 2523;  
Best Local Similarity 100.0%; Pred. No. 9.7e-18;  
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 292 TGCACCTTGGCGCTCTCTCGACCTTTACCTGGTCAAGGACGCGCGATGTCATTCC 347  
DB 1000 TGCACCTTGGCGCTCTCTCGACCTTTACCTGGTCAAGGACGCGCGATGTCATTCC 1055  
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Job time : 791 secs

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OM nucleic - nucleic search, using sw model

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6611.294 Million cell updates/sec

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Scoring table: OLIGO\_NUC  
Gapop 60.0 , Gapext 60.0

Searched: 682709 seqs, 277475446 residues

Word size : 35

Total number of hits satisfying chosen parameters: 116

Minimum DB seq length: 0  
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Post-processing: Listing first 1000 summaries

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6: /cgn2\_6/prodata/2/ina/backfiles1.seq.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	62	3.0	943	2	US-08-483-695-6
2	62	3.0	943	2	US-07-965-285-6
3	62	3.0	943	2	US-08-487-231-6
4	62	3.0	943	3	US-09-201-912-6
5	58	2.8	382	3	US-08-867-611-56
6	58	2.8	382	4	US-09-690-359-56
7	58	2.8	1414	3	US-08-867-611-51
8	58	2.8	1414	4	US-09-690-359-51
9	58	2.8	1420	3	US-08-867-611-57
10	58	2.8	1420	4	US-09-690-359-57
11	56	2.7	281	1	US-08-350-884-75
12	56	2.7	281	1	US-08-440-548-75
13	56	2.7	281	1	US-08-709-173-75
14	56	2.7	281	2	US-08-709-173-75
15	56	2.7	283	3	US-08-444-818-33
16	56	2.7	368	1	US-08-350-884-71
17	56	2.7	368	1	US-08-440-548-71
18	56	2.7	368	1	US-08-709-173-71
19	56	2.7	368	2	US-08-709-173-71
20	56	2.7	2058	4	US-09-881-654-1
21	56	2.7	2058	4	US-09-881-654-1
22	56	2.7	2064	1	US-08-350-884-69
23	56	2.7	2064	1	US-08-440-548-69
24	56	2.7	2064	1	US-08-709-173-69
25	56	2.7	2064	2	US-08-709-173-69
26	56	2.7	2523	1	US-08-350-884-85
27	56	2.7	2523	1	US-08-440-548-85

28	56	2.7	2523	1	US-08-709-173-85
29	56	2.7	2523	2	US-08-709-173-85
30	56	2.7	5360	3	US-08-444-818-53
31	56	2.7	5360	3	US-08-444-818-53
32	56	2.7	5360	3	US-08-444-818-53
33	56	2.7	7310	3	US-08-444-818-74
34	56	2.7	7310	3	US-08-444-818-74
35	56	2.7	8987	3	US-08-444-818-137
36	56	2.7	8987	3	US-08-444-818-137
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67	56	2.7	9185	3	US-08-444-818-122
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76	56	2.7	9185	3	US-08-444-818-122
77	56	2.7	9185	3	US-08-444-818-122
78	56	2.7	9185	3	US-08-444-818-122
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85	56	2.7	9185	3	US-08-444-818-122
86	56	2.7	9185	3	US-08-444-818-122
87	56	2.7	9185	3	US-08-444-818-122
88	56	2.7	9185	3	US-08-444-818-122
89	56	2.7	9185	3	US-08-444-818-122
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96	56	2.7	9185	3	US-08-444-818-122
97	56	2.7	9185	3	US-08-444-818-122
98	56	2.7	9185	3	US-08-444-818-122
99	56	2.7	9185	3	US-08-444-818-122
100	56	2.7	9185	3	US-08-444-818-122



RESULT 3  
US-08-487-231-6  
; Sequence 6, Application US/08487231  
; Patent No. 5919454  
; GENERAL INFORMATION:  
; APPLICANT: Brechot, Christian  
; APPLICANT: Kremsdorf, Dina  
; APPLICANT: Porchon, Colette  
; TITLE OF INVENTION: Nucleotide and Peptide Sequences of a  
; TITLE OF INVENTION: Hepatitis C Virus Isolate, Diagnostic and Therapeutic  
; TITLE OF INVENTION: Applications  
; NUMBER OF SEQUENCES: 46  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Finnegan, Henderson, Farabow, Garrett &  
; ADDRESSEE: Dunner  
; STREET: 1300 I Street, N.W.  
; CITY: Washington  
; STATE: DC  
; COUNTRY: USA  
; ZIP: 20005-3315  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patent In Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/487,231  
; FILING DATE: 07-JUNE-1995  
; CLASSIFICATION: 435  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: US 07/965,285  
; FILING DATE: 18-MAR-1993  
; CLASSIFICATION: 435  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: FR 91 06 882  
; FILING DATE: 06-JUN-1991  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Meyers, Kenneth J.  
; REGISTRATION NUMBER: 25,146  
; REFERENCE/DOCKET NUMBER: 05286-0001-02000  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: 202-408-4000  
; TELEFAX: 202-408-4400  
; INFORMATION FOR SEQ ID NO: 6:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 943 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; MOLECULE TYPE: Other  
; DESCRIPTION: cdna to genomic RNA  
US-08-487-231-6  
Query Match 3.0%; Score 62; DB 2; Length 943;  
Best Local Similarity 100.0%; Pred. No. 8.8e-22;  
Matches 62; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1915 GCGCGGCTTCGCTGCTTTGGCGCGGTATTGCTATCCACAGGCTGCGTGCATAGTA 1974  
Db 630 GCGCGGCTTCGCTGCTTTGGCGCGGTATTGCTATCCACAGGCTGCGTGCATAGTA 689  
QY 1975 GG 1976  
Db 690 GG 691  
RESULT 4  
US-09-201-912-6  
; Sequence 6, Application US/09201912  
; Patent No. 6210962  
; GENERAL INFORMATION:  
; APPLICANT: Brechot, Christian

APPLICANT: Kremsdorf, Dina  
APPLICANT: Porchon, Colette  
TITLE OF INVENTION: Nucleotide and Peptide Sequences of a  
TITLE OF INVENTION: Hepatitis C Virus Isolate, Diagnostic and Therapeutic  
TITLE OF INVENTION: Applications  
NUMBER OF SEQUENCES: 46  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Finnegan, Henderson, Farabow, Garrett &  
ADDRESSEE: Dunner  
STREET: 1300 I Street, N.W.  
CITY: Washington  
STATE: DC  
COUNTRY: USA  
ZIP: 20005-3315  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patent In Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/201,912  
FILING DATE:  
CLASSIFICATION:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 07/965,285  
FILING DATE:  
ATTORNEY/AGENT INFORMATION:  
NAME: Meyers, Kenneth J.  
REGISTRATION NUMBER: 25,146  
REFERENCE/DOCKET NUMBER: 05286-0001-00000  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 202-408-4000  
TELEFAX: 202-408-4400  
INFORMATION FOR SEQ ID NO: 6:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 943 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: Other  
DESCRIPTION: cdna to genomic RNA  
US-09-201-912-6  
Query Match 3.0%; Score 62; DB 3; Length 943;  
Best Local Similarity 100.0%; Pred. No. 8.8e-22;  
Matches 62; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1915 GCGCGGCTTCGCTGCTTTGGCGCGGTATTGCTATCCACAGGCTGCGTGCATAGTA 1974  
Db 630 GCGCGGCTTCGCTGCTTTGGCGCGGTATTGCTATCCACAGGCTGCGTGCATAGTA 689  
QY 1975 GG 1976  
Db 690 GG 691  
RESULT 5  
US-08-867-611-56  
; Sequence 36, Application US/08867611  
; Patent No. 6122189  
; GENERAL INFORMATION:  
; APPLICANT: DEVARE, SUSHIL G  
; APPLICANT: DESAI, SURESH M  
; APPLICANT: CASEY, JAMES M  
; APPLICANT: DAILEY, STEPHEN H  
; APPLICANT: DAWSON, GEORGE J  
; APPLICANT: GUTIERREZ, ROBIN A  
; APPLICANT: LESNIEWSKI, RICHARD R  
; APPLICANT: STEWART, JAMES L  
; APPLICANT: RUPPRECHT, KEVIN R  
; TITLE OF INVENTION: HEPATITIS C ASSAY UTILIZING RECOMBINANT  
; TITLE OF INVENTION: ANTIGENS  
; NUMBER OF SEQUENCES: 59





APPLICANT: DAWSON, GEORGE J  
APPLICANT: GUTIERREZ, ROBIN A  
APPLICANT: LESNIEWSKI, RICHARD R  
APPLICANT: STEWART, JAMES L  
APPLICANT: RUPPRECHT, KEVIN R  
TITLE OF INVENTION: HEPATITIS C ASSAY UTILIZING RECOMBINANT  
TITLE OF INVENTION: ANTIGENS  
NUMBER OF SEQUENCES: 59  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: ABBOTT LABORATORIES  
STREET: ONE ABBOTT PARK ROAD, CHAD377/AP6D2  
CITY: ABBOTT PARK  
STATE: IL  
COUNTRY: USA  
ZIP: 60064-3500  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patent In Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/867,611  
FILING DATE: 02-JUN-1997  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US/08/646,757  
FILING DATE:  
APPLICATION NUMBER: US/08/179,896  
FILING DATE:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/572,822  
FILING DATE: 24-AUG-1990  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/614,069  
FILING DATE: 07-NOV-1990  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/748,561  
FILING DATE: 21-AUG-1991  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/748,565  
FILING DATE: 21-AUG-1991  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/748,566  
FILING DATE: 21-AUG-1991  
ATTORNEY/AGENT INFORMATION:  
NAME: FOREMSKI, PRISCILLA E  
REGISTRATION NUMBER: 33,207  
REFERENCE/DOCKET NUMBER: 4834.US.P6  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 708-937-6365  
TELEFAX: 708-937-9556  
INFORMATION FOR SEQ ID NO: 51:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 1414 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
US-08-867-611-51  
Query Match 2.8%; Score 58; DB 3; Length 1414;  
Best Local Similarity 100.0%; Pred. No. 9.9e-20;  
Matches 58; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 1291 ACGTGTGTCACCCAGACAGTCGACTTCAGCCTTGACCTTACCTTACCATTTGAGACAA 1348  
Db 22 ACGTGTGTCACCCAGACAGTCGACTTCAGCCTTGACCTTACCTTACCATTTGAGACAA 79  
RESULT 8  
US-09-690-359-51  
; Sequence 51, Application US/09690359  
; Patent No. 6593083

GENERAL INFORMATION:  
APPLICANT: DEVARE, SUSHIL G  
APPLICANT: DESAI, SURESH M  
APPLICANT: CASEY, JAMES M  
APPLICANT: DAILEY, STEPHEN H  
APPLICANT: DAWSON, GEORGE J  
APPLICANT: GUTIERREZ, ROBIN A  
APPLICANT: LESNIEWSKI, RICHARD R  
APPLICANT: STEWART, JAMES L  
APPLICANT: RUPPRECHT, KEVIN R  
TITLE OF INVENTION: HEPATITIS C ASSAY UTILIZING RECOMBINANT  
TITLE OF INVENTION: ANTIGENS  
NUMBER OF SEQUENCES: 59  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: ABBOTT LABORATORIES  
STREET: ONE ABBOTT PARK ROAD, CHAD377/AP6D2  
CITY: ABBOTT PARK  
STATE: IL  
COUNTRY: USA  
ZIP: 60064-3500  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patent In Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/690,359  
FILING DATE: 17-Oct-2000  
CLASSIFICATION: <Unknown>  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/867,611  
FILING DATE: 02-JUN-1997  
APPLICATION NUMBER: US/08/646,757  
FILING DATE: <Unknown>  
APPLICATION NUMBER: US 07/572,822  
FILING DATE: <Unknown>  
APPLICATION NUMBER: US 07/614,069  
FILING DATE: 07-NOV-1990  
APPLICATION NUMBER: US 07/748,561  
FILING DATE: 21-AUG-1991  
APPLICATION NUMBER: US 07/748,565  
FILING DATE: 21-AUG-1991  
APPLICATION NUMBER: US 07/748,566  
FILING DATE: 21-AUG-1991  
ATTORNEY/AGENT INFORMATION:  
NAME: FOREMSKI, PRISCILLA E  
REGISTRATION NUMBER: 33,207  
REFERENCE/DOCKET NUMBER: 4834.US.P6  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 708-937-6365  
TELEFAX: 708-937-9556  
INFORMATION FOR SEQ ID NO: 51:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 1414 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
US-09-690-359-51  
Query Match 2.8%; Score 58; DB 4; Length 1414;  
Best Local Similarity 100.0%; Pred. No. 9.9e-20;  
Matches 58; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 1291 ACGTGTGTCACCCAGACAGTCGACTTCAGCCTTGACCTTACCTTACCATTTGAGACAA 1348  
Db 22 ACGTGTGTCACCCAGACAGTCGACTTCAGCCTTGACCTTACCTTACCATTTGAGACAA 79  
RESULT 9





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; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/680,296
; FILING DATE: 04-APR-1991
; ATTORNEY/AGENT INFORMATION:
; NAME: CIOTTI, THOMAS E.
; REGISTRATION NUMBER: 21,013
; REFERENCE/DOCKET NUMBER: 22300-20100.20
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 813-5600
; TELEFAX: (415) 494-0792
; TELEX: 706141
; INFORMATION FOR SEQ ID NO: 75:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 281 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 1..279
; US-08-709-173-75

Query Match      2.7%; Score 56; DB 1; Length 281;
Best Local Similarity 100.0%; Pred. No. 1.1e-18;
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      292 TGCACCTTGGCGCTCTCTCGGACCTTTACCTGCTCAGGAGGACGCCGATGTCATTCC 347
Db      4 TGCACCTTGGCGCTCTCTCGGACCTTTACCTGCTCAGGAGGACGCCGATGTCATTCC 59

RESULT 14
US-08-709-177-75
; Sequence 75, Application US/08709177
; Patent No. 5885799
; GENERAL INFORMATION:
; APPLICANT: HOUGHTON, MICHAEL
; APPLICANT: CHOO, QUI LIM
; TITLE OF INVENTION: HEPATITIS C VIRUS PROTEASE
; NUMBER OF SEQUENCES: 86
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: MORRISON & FOERSTER
; STREET: 755 Page Mill Road
; City: Palo Alto
; STATE: California
; COUNTRY: USA
; ZIP: 94304-1018
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent in Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/709,177
; FILING DATE: 06-SEP-1996
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/680,296
; FILING DATE: 04-APR-1991
; ATTORNEY/AGENT INFORMATION:
; NAME: CIOTTI, THOMAS E.
; REGISTRATION NUMBER: 21,013
; REFERENCE/DOCKET NUMBER: 22300-20100.20
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 813-5600
; TELEFAX: (415) 494-0792
; TELEX: 706141
; INFORMATION FOR SEQ ID NO: 75:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 281 base pairs
; TYPE: nucleic acid
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; STRANDEDNESS: single
; TOPOLOGY: linear
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 1..279
; US-08-709-177-75

Query Match      2.7%; Score 56; DB 2; Length 281;
Best Local Similarity 100.0%; Pred. No. 1.1e-18;
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      292 TGCACCTTGGCGCTCTCTCGGACCTTTACCTGCTCAGGAGGACGCCGATGTCATTCC 347
Db      4 TGCACCTTGGCGCTCTCTCGGACCTTTACCTGCTCAGGAGGACGCCGATGTCATTCC 59

RESULT 15
US-08-444-818-33
; Sequence 33, Application US/08444818
; Patent No. 6150087
; GENERAL INFORMATION:
; APPLICANT: Chien, David Y.
; APPLICANT: Rutter, William J.
; TITLE OF INVENTION: NANV Diagnostics and Vaccines
; NUMBER OF SEQUENCES: 777
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Chiron Corporation
; STREET: 4560 Horton Street
; City: Emeryville
; STATE: CA
; COUNTRY: USA
; ZIP: 94608-2916
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent in Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/444,818
; FILING DATE:
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/08/403,590
; FILING DATE: 14-MAR-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Harbin, Alisa A.
; REGISTRATION NUMBER: 33,895
; REFERENCE/DOCKET NUMBER: 0110.002
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (508)359-3876
; TELEFAX: (508)359-3885
; INFORMATION FOR SEQ ID NO: 33:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 283 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: CDNA
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 3..281
; US-08-444-818-33

Query Match      2.7%; Score 56; DB 3; Length 283;
Best Local Similarity 100.0%; Pred. No. 1.1e-18;
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      292 TGCACCTTGGCGCTCTCTCGGACCTTTACCTGCTCAGGAGGACGCCGATGTCATTCC 347
Db      6 TGCACCTTGGCGCTCTCTCGGACCTTTACCTGCTCAGGAGGACGCCGATGTCATTCC 61

RESULT 16
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US-08-350-884-71  
; Sequence 71 Application US/08350884  
; Patent No. 5597591

## GENERAL INFORMATION:

APPLICANT: HOUGHTON, MICHAEL  
APPLICANT: CHOO, QUI LIM  
APPLICANT: KOO, GEORGE  
TITLE OF INVENTION: HEPATITIS C VIRUS PROTEASE  
NUMBER OF SEQUENCES: 86  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: MORRISON & FOERSTER  
STREET: 755 Page Mill Road  
CITY: Palo Alto  
STATE: California  
COUNTRY: USA  
ZIP: 94304-1018

## COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patent In Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/350,884  
FILING DATE: 06-DEC-1994  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/680,296  
FILING DATE: 04-APR-1991  
ATTORNEY/AGENT INFORMATION:  
NAME: CIOTTI, THOMAS E.  
REGISTRATION NUMBER: 21,013  
REFERENCE/DOCKET NUMBER: 22300-20100.20  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (415) 813-5600  
TELEFAX: (415) 494-0792

## INFORMATION FOR SEQ ID NO: 71:

SEQUENCE CHARACTERISTICS:  
LENGTH: 368 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
FEATURE:  
NAME/KEY: CDS  
LOCATION: 1..366  
US-08-350-884-71

Query Match 2.7%; Score 56; DB 1; Length 368;  
Best Local Similarity 100.0%; Pred. No. 1.1e-18;  
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 292 TGCACCTTGGCGCTCTCGGACCTTTACTGTGTCACGAGGACGCGGATGTCATTC 347  
Db 220 TGCACCTTGGCGCTCTCGGACCTTTACTGTGTCACGAGGACGCGGATGTCATTC 275

## RESULT 17

US-08-440-548-71  
; Sequence 71 Application US/08440548  
; Patent No. 5597591

## GENERAL INFORMATION:

APPLICANT: HOUGHTON, MICHAEL  
APPLICANT: CHOO, QUI LIM  
APPLICANT: KOO, GEORGE  
TITLE OF INVENTION: HEPATITIS C VIRUS PROTEASE  
NUMBER OF SEQUENCES: 86  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: MORRISON & FOERSTER  
STREET: 755 Page Mill Road  
CITY: Palo Alto  
STATE: California  
COUNTRY: USA  
ZIP: 94304-1018

## COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patent In Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/440,548  
FILING DATE: 12-MAY-1995  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/680,296  
FILING DATE: 04-APR-1991  
ATTORNEY/AGENT INFORMATION:  
NAME: CIOTTI, THOMAS E.  
REGISTRATION NUMBER: 21,013  
REFERENCE/DOCKET NUMBER: 22300-20100.20  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (415) 813-5600  
TELEFAX: (415) 494-0792  
TELEX: 706141

## INFORMATION FOR SEQ ID NO: 71:

SEQUENCE CHARACTERISTICS:  
LENGTH: 368 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
FEATURE:  
NAME/KEY: CDS  
LOCATION: 1..366  
US-08-440-548-71

Query Match 2.7%; Score 56; DB 1; Length 368;  
Best Local Similarity 100.0%; Pred. No. 1.1e-18;  
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 292 TGCACCTTGGCGCTCTCGGACCTTTACTGTGTCACGAGGACGCGGATGTCATTC 347  
Db 220 TGCACCTTGGCGCTCTCGGACCTTTACTGTGTCACGAGGACGCGGATGTCATTC 275

## RESULT 18

US-08-709-125-71  
; Sequence 71 Application US/08709173  
; Patent No. 5712145

## GENERAL INFORMATION:

APPLICANT: HOUGHTON, MICHAEL  
APPLICANT: CHOO, QUI LIM  
APPLICANT: KOO, GEORGE  
TITLE OF INVENTION: HEPATITIS C VIRUS PROTEASE  
NUMBER OF SEQUENCES: 86  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: MORRISON & FOERSTER  
STREET: 755 Page Mill Road  
CITY: Palo Alto  
STATE: California  
COUNTRY: USA  
ZIP: 94304-1018

## COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patent In Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/709,173  
FILING DATE: 06-SEP-1996  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/680,296  
FILING DATE: 04-APR-1991  
ATTORNEY/AGENT INFORMATION:  
NAME: CIOTTI, THOMAS E.  
REGISTRATION NUMBER: 21,013  
REFERENCE/DOCKET NUMBER: 22300-20100.20

TELECOMMUNICATION INFORMATION:  
TELEPHONE: (415) 813-5600  
TELEFAX: (415) 494-0792  
TELEX: 706141  
INFORMATION FOR SEQ ID NO: 71:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 368 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
FEATURE:  
NAME/KEY: CDS  
LOCATION: 1..366  
US-08-709-173-71

Query Match 2.7%; Score 56; DB 1; Length 368;  
Best Local Similarity 100.0%; Pred. No. 1.1e-18;  
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 292 TGCACCTTGGCGCTCTCGGACCTTTACTGTCACGAGGACGCCGATGTCATTCC 347  
Db 220 TGCACCTTGGCGCTCTCGGACCTTTACTGTCACGAGGACGCCGATGTCATTCC 275

RESULT 19  
US-08-709-177-71  
Sequence 71 Application US/08709177  
Patent No. 6632601  
GENERAL INFORMATION:  
APPLICANT: HOUGHTON, MICHAEL  
APPLICANT: CHOO, QUI LIM  
APPLICANT: KUO, GEORGE  
TITLE OF INVENTION: HEPATITIS C VIRUS PROTEASE  
NUMBER OF SEQUENCES: 86  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: MORRISON & FOERSTER  
STREET: 755 Page Mill Road  
CITY: Palo Alto  
STATE: California  
COUNTRY: USA  
ZIP: 94304-1018  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/709,177  
FILING DATE: 06-SEP-1996  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/680,296  
FILING DATE: 04-APR-1991  
ATTORNEY/AGENT INFORMATION:  
NAME: CIOTTI, THOMAS E.  
REGISTRATION NUMBER: 21,013  
REFERENCE/DOCKET NUMBER: 22300-20100.20  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (415) 813-5600  
TELEFAX: (415) 494-0792  
TELEX: 706141  
INFORMATION FOR SEQ ID NO: 71:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 368 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
FEATURE:  
NAME/KEY: CDS  
LOCATION: 1..366  
US-08-709-177-71

Query Match 2.7%; Score 56; DB 2; Length 368;

Best Local Similarity 100.0%; Pred. No. 1.1e-18;  
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 292 TGCACCTTGGCGCTCTCGGACCTTTACTGTCACGAGGACGCCGATGTCATTCC 347  
Db 220 TGCACCTTGGCGCTCTCGGACCTTTACTGTCACGAGGACGCCGATGTCATTCC 275

RESULT 20  
US-09-881-239-2  
Sequence 2 Application US/09881239  
Patent No. 6630298  
GENERAL INFORMATION:  
APPLICANT: CHIEN, David Y.  
APPLICANT: ARCANGEL, Phillip  
APPLICANT: TANDESKE, Laura  
APPLICANT: GEORGE-NASCIEMENTO, Carlos  
APPLICANT: COIT, Doris  
APPLICANT: MEDINA-SELBY, Angelica  
TITLE OF INVENTION: HCV ANTIGEN/ANTIBODY COMBINATION ASSAY  
FILE REFERENCE: 2302-16073 / PPI6073.003  
CURRENT APPLICATION NUMBER: US/09/881,239  
CURRENT FILING DATE: 2001-06-14  
NUMBER OF SEQ ID NOS: 8  
SOFTWARE: PatentIn Ver. 2.0  
SEQ ID NO 2  
LENGTH: 2058  
TYPE: DNA  
ORGANISM: Artificial Sequence  
FEATURE:  
OTHER INFORMATION: Description of Artificial Sequence:  
OTHER INFORMATION: representative NS3/4a conformational antigen  
NAME/KEY: CDS  
LOCATION: (1)..(2058)  
US-09-881-239-2

Query Match 2.7%; Score 56; DB 4; Length 2058;  
Best Local Similarity 100.0%; Pred. No. 1e-18;  
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 292 TGCACCTTGGCGCTCTCGGACCTTTACTGTCACGAGGACGCCGATGTCATTCC 347  
Db 292 TGCACCTTGGCGCTCTCGGACCTTTACTGTCACGAGGACGCCGATGTCATTCC 347

RESULT 21  
US-09-881-654-1  
Sequence 1 Application US/09881654  
Patent No. 6632601  
GENERAL INFORMATION:  
APPLICANT: CHIEN, David Y.  
APPLICANT: ARCANGEL, Phillip  
APPLICANT: TANDESKE, Laura  
APPLICANT: GEORGE-NASCIEMENTO, Carlos  
APPLICANT: COIT, Doris  
APPLICANT: MEDINA-SELBY, Angelica  
TITLE OF INVENTION: IMMUNOASSAYS FOR ANTI-HCV ANTIBODIES  
FILE REFERENCE: 2302-17039 / PPI7039.002  
CURRENT APPLICATION NUMBER: US/09/881,654  
CURRENT FILING DATE: 2001-06-14  
PRIOR APPLICATION NUMBER: 60/212,082  
PRIOR FILING DATE: 2000-06-15  
PRIOR APPLICATION NUMBER: 60/280,811  
PRIOR FILING DATE: 2001-04-02  
PRIOR APPLICATION NUMBER: 60/280,867  
PRIOR FILING DATE: 2001-04-02  
NUMBER OF SEQ ID NOS: 7  
SOFTWARE: PatentIn Ver. 2.0  
SEQ ID NO 1  
LENGTH: 2058  
TYPE: DNA  
ORGANISM: Artificial Sequence  
FEATURE:

; OTHER INFORMATION: Description of Artificial Sequence:  
; OTHER INFORMATION: representative N33/4a conformational antigen  
; NAME/KEY: CDS  
; LOCATION: (1)..(2058)  
US-09-881-654-1

Query Match 2.7%; Score 56; DB 4; Length 2058;  
Best Local Similarity 100.0%; Pred. No. 1e-18;  
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTGGGCTCTCGGACCTTTACCTGGTACAGGAGCAGCCGATGTCATTC 347  
DB 292 TGCACCTTGGGCTCTCGGACCTTTACCTGGTACAGGAGCAGCCGATGTCATTC 347

RESULT 22

US-08-350-884-69  
; Sequence 69, Application US/08350884  
; Patent No. 5585258  
; GENERAL INFORMATION:  
; APPLICANT: HOUGHTON, MICHAEL  
; APPLICANT: CHOO, QUI LIM  
; APPLICANT: KUI, GEORGE  
; TITLE OF INVENTION: HEPATITIS C VIRUS PROTEASE  
; NUMBER OF SEQUENCES: 86  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: MORRISON & FOERSTER  
; STREET: 755 Page Mill Road  
; CITY: Palo Alto  
; STATE: California  
; COUNTRY: USA  
; ZIP: 94304-1018  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patent In Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/350,884  
; FILING DATE: 06-DEC-1994  
; CLASSIFICATION: 435  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: US 07/680,296  
; FILING DATE: 04-APR-1991  
; ATTORNEY/AGENT INFORMATION:  
; NAME: CIOTTI, THOMAS E.  
; REGISTRATION NUMBER: 21,013  
; REFERENCE/DOCKET NUMBER: 22300-20100.20  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (415) 813-5600  
; TELEFAX: (415) 494-0792  
; TELEX: 706141  
; INFORMATION FOR SEQ ID NO: 69:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 2064 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; FEATURE:  
; NAME/KEY: CDS  
; LOCATION: 7..2064  
US-08-350-884-69

Query Match 2.7%; Score 56; DB 1; Length 2064;  
Best Local Similarity 100.0%; Pred. No. 1e-18;  
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTGGGCTCTCGGACCTTTACCTGGTACAGGAGCAGCCGATGTCATTC 347  
DB 541 TGCACCTTGGGCTCTCGGACCTTTACCTGGTACAGGAGCAGCCGATGTCATTC 596

RESULT 23

US-08-440-548-69  
; Sequence 69, Application US/08440548  
; Patent No. 5597691  
; GENERAL INFORMATION:  
; APPLICANT: HOUGHTON, MICHAEL  
; APPLICANT: CHOO, QUI LIM  
; APPLICANT: KUI, GEORGE  
; TITLE OF INVENTION: HEPATITIS C VIRUS PROTEASE  
; NUMBER OF SEQUENCES: 86  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: MORRISON & FOERSTER  
; STREET: 755 Page Mill Road  
; CITY: Palo Alto  
; STATE: California  
; COUNTRY: USA  
; ZIP: 94304-1018  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patent In Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/440,548  
; FILING DATE: 12-MAY-1995  
; CLASSIFICATION: 435  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: US 07/680,296  
; FILING DATE: 04-APR-1991  
; ATTORNEY/AGENT INFORMATION:  
; NAME: CIOTTI, THOMAS E.  
; REGISTRATION NUMBER: 21,013  
; REFERENCE/DOCKET NUMBER: 22300-20100.20  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (415) 813-5600  
; TELEFAX: (415) 494-0792  
; TELEX: 706141  
; INFORMATION FOR SEQ ID NO: 69:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 2064 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; FEATURE:  
; NAME/KEY: CDS  
; LOCATION: 7..2064  
US-08-440-548-69

Query Match 2.7%; Score 56; DB 1; Length 2064;  
Best Local Similarity 100.0%; Pred. No. 1e-18;  
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTGGGCTCTCGGACCTTTACCTGGTACAGGAGCAGCCGATGTCATTC 347  
DB 541 TGCACCTTGGGCTCTCGGACCTTTACCTGGTACAGGAGCAGCCGATGTCATTC 596

RESULT 24

US-08-709-173-69  
; Sequence 69, Application US/08709173  
; Patent No. 5712145  
; GENERAL INFORMATION:  
; APPLICANT: HOUGHTON, MICHAEL  
; APPLICANT: CHOO, QUI LIM  
; APPLICANT: KUI, GEORGE  
; TITLE OF INVENTION: HEPATITIS C VIRUS PROTEASE  
; NUMBER OF SEQUENCES: 86  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: MORRISON & FOERSTER  
; STREET: 755 Page Mill Road  
; CITY: Palo Alto  
; STATE: California  
; COUNTRY: USA  
; ZIP: 94304-1018

COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/709,173  
FILING DATE: 06-SEP-1996  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/680,296  
FILING DATE: 04-APR-1991  
ATTORNEY/AGENT INFORMATION:  
NAME: CIOTTI, THOMAS E.  
REGISTRATION NUMBER: 21,013  
REFERENCE/DOCKET NUMBER: 22300-20100.20  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (415) 813-5600  
TELEFAX: (415) 494-0792  
TELEX: 706141  
INFORMATION FOR SEQ ID NO: 69:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 2064 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
FEATURE:  
NAME/KEY: CDS  
LOCATION: 7..2064  
US-08-709-173-69

Query Match 2.7%; Score 56; DB 1; Length 2064;  
Best Local Similarity 100.0%; Pred. No. 1e-18;  
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTGGCGCTCTCGGACCTTTACTGTCACGAGGCACGCCGATGTCATTC 347  
DB 541 TGCACCTTGGCGCTCTCGGACCTTTACTGTCACGAGGCACGCCGATGTCATTC 596

RESULT 25  
US-08-709-173-69  
Sequence 69, Application US/08/709177  
Patent No. 5885799  
GENERAL INFORMATION:  
APPLICANT: HOUGHTON, MICHAEL  
APPLICANT: CHOO, QUI LIM  
TITLE OF INVENTION: HEPATITIS C VIRUS PROTEASE  
NUMBER OF SEQUENCES: 86  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: MORRISON & FOERSTER  
STREET: 755 Page Mill Road  
CITY: Palo Alto  
STATE: California  
COUNTRY: USA  
ZIP: 94304-1018  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/709,177  
FILING DATE: 06-SEP-1996  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/680,296  
FILING DATE: 04-APR-1991  
ATTORNEY/AGENT INFORMATION:  
NAME: CIOTTI, THOMAS E.  
REGISTRATION NUMBER: 21,013  
REFERENCE/DOCKET NUMBER: 22300-20100.20

TELECOMMUNICATION INFORMATION:  
TELEPHONE: (415) 813-5600  
TELEFAX: (415) 494-0792  
TELEX: 706141  
INFORMATION FOR SEQ ID NO: 69:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 2064 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
FEATURE:  
NAME/KEY: CDS  
LOCATION: 7..2064  
US-08-709-177-69

Query Match 2.7%; Score 56; DB 2; Length 2064;  
Best Local Similarity 100.0%; Pred. No. 1e-18;  
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTGGCGCTCTCGGACCTTTACTGTCACGAGGCACGCCGATGTCATTC 347  
DB 541 TGCACCTTGGCGCTCTCGGACCTTTACTGTCACGAGGCACGCCGATGTCATTC 596

RESULT 26  
US-08-350-884-85  
Sequence 85, Application US/08350884  
Patent No. 5585258  
GENERAL INFORMATION:  
APPLICANT: HOUGHTON, MICHAEL  
APPLICANT: CHOO, QUI LIM  
TITLE OF INVENTION: HEPATITIS C VIRUS PROTEASE  
NUMBER OF SEQUENCES: 86  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: MORRISON & FOERSTER  
STREET: 755 Page Mill Road  
CITY: Palo Alto  
STATE: California  
COUNTRY: USA  
ZIP: 94304-1018  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/350,884  
FILING DATE: 06-DEC-1994  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/680,296  
FILING DATE: 04-APR-1991  
ATTORNEY/AGENT INFORMATION:  
NAME: CIOTTI, THOMAS E.  
REGISTRATION NUMBER: 21,013  
REFERENCE/DOCKET NUMBER: 22300-20100.20  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (415) 813-5600  
TELEFAX: (415) 494-0792  
TELEX: 706141  
INFORMATION FOR SEQ ID NO: 85:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 2523 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
FEATURE:  
NAME/KEY: CDS  
LOCATION: 1..2523  
US-08-350-884-85

Query Match 2.7%; Score 56; DB 1; Length 2523;



Best Local Similarity 100.0%; Pred. No. 1e-18; Mismatches 0; Indels 0; Gaps 0;  
Matches 56; Conservative 0;

QY 292 TGCACCTGGGCTCTCGGACCTTACCTGGTCAAGGACGCGCGATGTCATTC 347  
DB 1000 TGCACCTGGGCTCTCGGACCTTACCTGGTCAAGGACGCGCGATGTCATTC 1055

RESULT 27  
US-08-440-548-85  
; Sequence 85, Application US/08440548  
; Patent No. 5597691  
; GENERAL INFORMATION:  
; APPLICANT: HOUGHTON, MICHAEL  
; APPLICANT: CHOO, QUI LIM  
; APPLICANT: KUI, GEORGE  
; TITLE OF INVENTION: HEPATITIS C VIRUS PROTEASE  
; NUMBER OF SEQUENCES: 86  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: MORRISON & FOERSTER  
; STREET: 755 Page Mill Road  
; CITY: Palo Alto  
; STATE: California  
; COUNTRY: USA  
; ZIP: 94304-1018  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patent In Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/440,548  
; FILING DATE: 12-MAY-1995  
; CLASSIFICATION: 435  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: US 07/680,296  
; FILING DATE: 04-APR-1991  
; ATTORNEY/AGENT INFORMATION:  
; NAME: CIOTTI, THOMAS E.  
; REGISTRATION NUMBER: 21,013  
; REFERENCE/DOCKET NUMBER: 22300-20100.20  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (415) 813-5600  
; TELEFAX: (415) 494-0792  
; TELEX: 706141  
; INFORMATION FOR SEQ ID NO: 85:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 2523 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; FEATURE:  
; NAME/KEY: CDS  
; LOCATION: 1..2523  
; ATTORNEY/AGENT INFORMATION:  
; NAME: CIOTTI, THOMAS E.  
; REGISTRATION NUMBER: 21,013  
; REFERENCE/DOCKET NUMBER: 22300-20100.20  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (415) 813-5600  
; TELEFAX: (415) 494-0792  
; TELEX: 706141  
; INFORMATION FOR SEQ ID NO: 85:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 2523 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; FEATURE:  
; NAME/KEY: CDS  
; LOCATION: 1..2523  
US-08-440-548-85

Query Match 2.7%; Score 56; DB 1; Length 2523;  
Best Local Similarity 100.0%; Pred. No. 1e-18;  
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTGGGCTCTCGGACCTTACCTGGTCAAGGACGCGCGATGTCATTC 347  
DB 1000 TGCACCTGGGCTCTCGGACCTTACCTGGTCAAGGACGCGCGATGTCATTC 1055

RESULT 28  
US-08-709-173-85  
; Sequence 85, Application US/08709173  
; Patent No. 5712145  
; GENERAL INFORMATION:  
; APPLICANT: HOUGHTON, MICHAEL  
; APPLICANT: CHOO, QUI LIM  
; APPLICANT: KUI, GEORGE  
; TITLE OF INVENTION: HEPATITIS C VIRUS PROTEASE  
; NUMBER OF SEQUENCES: 86  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: MORRISON & FOERSTER  
; STREET: 755 Page Mill Road  
; CITY: Palo Alto  
; STATE: California  
; COUNTRY: USA  
; ZIP: 94304-1018  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patent In Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/709,173  
; FILING DATE: 06-SEP-1996  
; CLASSIFICATION: 435  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: US 07/680,296  
; FILING DATE: 04-APR-1991  
; ATTORNEY/AGENT INFORMATION:  
; NAME: CIOTTI, THOMAS E.  
; REGISTRATION NUMBER: 21,013  
; REFERENCE/DOCKET NUMBER: 22300-20100.20  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (415) 813-5600  
; TELEFAX: (415) 494-0792  
; TELEX: 706141  
; INFORMATION FOR SEQ ID NO: 85:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 2523 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; FEATURE:  
; NAME/KEY: CDS  
; LOCATION: 1..2523  
US-08-709-173-85

Query Match 2.7%; Score 56; DB 1; Length 2523;  
Best Local Similarity 100.0%; Pred. No. 1e-18;  
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTGGGCTCTCGGACCTTACCTGGTCAAGGACGCGCGATGTCATTC 347  
DB 1000 TGCACCTGGGCTCTCGGACCTTACCTGGTCAAGGACGCGCGATGTCATTC 1055

RESULT 29  
US-08-709-177-85  
; Sequence 85, Application US/08709177  
; Patent No. 585799  
; GENERAL INFORMATION:  
; APPLICANT: HOUGHTON, MICHAEL  
; APPLICANT: CHOO, QUI LIM  
; APPLICANT: KUI, GEORGE  
; TITLE OF INVENTION: HEPATITIS C VIRUS PROTEASE  
; NUMBER OF SEQUENCES: 86  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: MORRISON & FOERSTER  
; STREET: 755 Page Mill Road  
; CITY: Palo Alto  
; STATE: California  
; COUNTRY: USA  
; ZIP: 94304-1018  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: Patent In Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/709,177

/ FILING DATE: 06-SEP-1996  
/ CLASSIFICATION: 435  
/ PRIOR APPLICATION DATA:  
/ APPLICATION NUMBER: US 07/680,296  
/ FILING DATE: 04-APR-1991  
/ ATTORNEY/AGENT INFORMATION:  
/ NAME: CIOTTI, THOMAS E.  
/ REGISTRATION NUMBER: 21,013  
/ REFERENCE/DOCKET NUMBER: 22300-20100.20  
/ TELECOMMUNICATION INFORMATION:  
/ TELEPHONE: (415) 813-5600  
/ TELEFAX: (415) 494-0792  
/ TELEX: 706141  
/ INFORMATION FOR SEQ ID NO: 85:  
/ SEQUENCE CHARACTERISTICS:  
/ LENGTH: 2523 base pairs  
/ TYPE: nucleic acid  
/ STRANDEDNESS: single  
/ TOPOLOGY: linear  
/ FEATURE:  
/ NAME/KEY: CDS  
/ LOCATION: 1..2523  
/ US-08-709-177-85

Query Match 2.7%; Score 56; DB 2; Length 2523;  
Best Local Similarity 100.0%; Pred. No. 1e-18;  
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTGGCGCTCCTCGGACCTTTACCTGTCACGAGGACGCCGATGTCATTCC 347  
Db 1000 TGCACCTTGGCGCTCCTCGGACCTTTACCTGTCACGAGGACGCCGATGTCATTCC 1055

RESULT 30  
US-08-444-818-53  
/ Sequence 53; Application US/08444818  
/ Patent No. 6150087  
/ GENERAL INFORMATION:  
/ APPLICANT: Chien, David Y.  
/ APPLICANT: Rutter, William J.  
/ TITLE OF INVENTION: NANBV Diagnostics and Vaccines  
/ NUMBER OF SEQUENCES: 777  
/ CORRESPONDENCE ADDRESS:  
/ ADDRESSEE: Chiron Corporation  
/ STREET: 4560 Horton Street  
/ CITY: Emeryville  
/ STATE: CA  
/ COUNTRY: USA  
/ ZIP: 94608-2916  
/ COMPUTER READABLE FORM:  
/ MEDIUM TYPE: Floppy disk  
/ COMPUTER: IBM PC compatible  
/ OPERATING SYSTEM: PC-DOS/MS-DOS  
/ SOFTWARE: Patent in Release #1.0, Version #1.30  
/ CURRENT APPLICATION DATA:  
/ FILING DATE: US/08/444,818  
/ PRIOR APPLICATION DATA:  
/ APPLICATION NUMBER: US/08/403,590  
/ FILING DATE: 14-MAR-1995  
/ ATTORNEY/AGENT INFORMATION:  
/ NAME: Harbin, Alisa A.  
/ REGISTRATION NUMBER: 33,895  
/ REFERENCE/DOCKET NUMBER: 0110.002  
/ TELECOMMUNICATION INFORMATION:  
/ TELEPHONE: (508) 359-3876  
/ TELEFAX: (508) 359-3885  
/ INFORMATION FOR SEQ ID NO: 65:  
/ SEQUENCE CHARACTERISTICS:  
/ LENGTH: 6785 base pairs  
/ TYPE: nucleic acid  
/ STRANDEDNESS: single  
/ TOPOLOGY: linear  
/ MEDIUM TYPE: CDNA  
/ FEATURE:  
/ NAME/KEY: CDS  
/ LOCATION: 3..6785  
/ US-08-444-818-65

Query Match 2.7%; Score 56; DB 3; Length 6785;  
Best Local Similarity 100.0%; Pred. No. 1e-18;  
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTGGCGCTCCTCGGACCTTTACCTGTCACGAGGACGCCGATGTCATTCC 347  
Db 1494 TGCACCTTGGCGCTCCTCGGACCTTTACCTGTCACGAGGACGCCGATGTCATTCC 1549

RESULT 32

/ TOPOLOGY: linear  
/ MOLECULE TYPE: CDNA  
/ FEATURE:  
/ NAME/KEY: CDS  
/ LOCATION: 3..5360  
/ US-08-444-818-53

Query Match 2.7%; Score 56; DB 3; Length 5360;  
Best Local Similarity 100.0%; Pred. No. 1e-18;  
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTGGCGCTCCTCGGACCTTTACCTGTCACGAGGACGCCGATGTCATTCC 347  
Db 1221 TGCACCTTGGCGCTCCTCGGACCTTTACCTGTCACGAGGACGCCGATGTCATTCC 1276

RESULT 31  
US-08-444-818-65  
/ Sequence 65; Application US/08444818  
/ Patent No. 6150087  
/ GENERAL INFORMATION:  
/ APPLICANT: Chien, David Y.  
/ APPLICANT: Rutter, William J.  
/ TITLE OF INVENTION: NANBV Diagnostics and Vaccines  
/ NUMBER OF SEQUENCES: 777  
/ CORRESPONDENCE ADDRESS:  
/ ADDRESSEE: Chiron Corporation  
/ STREET: 4560 Horton Street  
/ CITY: Emeryville  
/ STATE: CA  
/ COUNTRY: USA  
/ ZIP: 94608-2916  
/ COMPUTER READABLE FORM:  
/ MEDIUM TYPE: Floppy disk  
/ COMPUTER: IBM PC compatible  
/ OPERATING SYSTEM: PC-DOS/MS-DOS  
/ SOFTWARE: Patent in Release #1.0, Version #1.30  
/ CURRENT APPLICATION DATA:  
/ FILING DATE: US/08/444,818  
/ PRIOR APPLICATION DATA:  
/ APPLICATION NUMBER: US/08/403,590  
/ FILING DATE: 14-MAR-1995  
/ ATTORNEY/AGENT INFORMATION:  
/ NAME: Harbin, Alisa A.  
/ REGISTRATION NUMBER: 33,895  
/ REFERENCE/DOCKET NUMBER: 0110.002  
/ TELECOMMUNICATION INFORMATION:  
/ TELEPHONE: (508) 359-3876  
/ TELEFAX: (508) 359-3885  
/ INFORMATION FOR SEQ ID NO: 65:  
/ SEQUENCE CHARACTERISTICS:  
/ LENGTH: 6785 base pairs  
/ TYPE: nucleic acid  
/ STRANDEDNESS: single  
/ TOPOLOGY: linear  
/ MOLECULE TYPE: CDNA  
/ FEATURE:  
/ NAME/KEY: CDS  
/ LOCATION: 3..6785  
/ US-08-444-818-65

Query Match 2.7%; Score 56; DB 3; Length 6785;  
Best Local Similarity 100.0%; Pred. No. 1e-18;  
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 292 TGCACCTTGGCGCTCCTCGGACCTTTACCTGTCACGAGGACGCCGATGTCATTCC 347  
Db 1494 TGCACCTTGGCGCTCCTCGGACCTTTACCTGTCACGAGGACGCCGATGTCATTCC 1549

RESULT 32

24

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US-08-444-818-74
; Sequence 74 Application, US/08444818
; Patent No. 6150087
; GENERAL INFORMATION:
; APPLICANT: Chien, David Y.
; APPLICANT: Rutter, William J.
; TITLE OF INVENTION: NANBV Diagnostics and Vaccines
; NUMBER OF SEQUENCES: 777
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Chiron Corporation
; STREET: 4560 Horton Street
; CITY: Emeryville
; STATE: CA
; COUNTRY: USA
; ZIP: 94608-2916
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/444,818
; FILING DATE:
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/08/403,590
; FILING DATE: 14-MAR-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Harbin, Alisa A.
; REGISTRATION NUMBER: 33,895
; REFERENCE/DOCKET NUMBER: 0110.002
; TELEPHONE: (508)359-3876
; TELEFAX: (508)359-3885
; INFORMATION FOR SEQ ID NO: 74:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 7310 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 3..7310
US-08-444-818-74

Query Match 2.7%; Score 56; DB 3; Length 7310;
Best Local Similarity 100.0%; Pred. No. 1e-18;
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 292 TGCACCTTGGCGCTCTCGGACCTTACCTGGTCACGAGGACCGCGATGTCATCC 347
Db 2019 TGCACCTTGGCGCTCTCGGACCTTACCTGGTCACGAGGACCGCGATGTCATCC 2074

RESULT 33
US-08-444-818-88
; Sequence 88 Application US/08444818
; Patent No. 6150087
; GENERAL INFORMATION:
; APPLICANT: Chien, David Y.
; APPLICANT: Rutter, William J.
; TITLE OF INVENTION: NANBV Diagnostics and Vaccines
; NUMBER OF SEQUENCES: 777
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Chiron Corporation
; STREET: 4560 Horton Street
; CITY: Emeryville
; STATE: CA
; COUNTRY: USA
; ZIP: 94608-2916
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/444,818
; FILING DATE:
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/08/403,590
; FILING DATE: 14-MAR-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Harbin, Alisa A.
; REGISTRATION NUMBER: 33,895
; REFERENCE/DOCKET NUMBER: 0110.002
; TELEPHONE: (508)359-3876
; TELEFAX: (508)359-3885

```

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COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent In Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/444,818
FILING DATE:
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/08/403,590
FILING DATE: 14-MAR-1995
ATTORNEY/AGENT INFORMATION:
NAME: Harbin, Alisa A.
REGISTRATION NUMBER: 33,895
REFERENCE/DOCKET NUMBER: 0110.002
TELEPHONE: (508)359-3876
TELEFAX: (508)359-3885
INFORMATION FOR SEQ ID NO: 88:
SEQUENCE CHARACTERISTICS:
LENGTH: 8316 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
FEATURE:
NAME/KEY: CDS
LOCATION: 1..8316
US-08-444-818-88

Query Match 2.7%; Score 56; DB 3; Length 8316;
Best Local Similarity 100.0%; Pred. No. 1e-18;
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 292 TGCACCTTGGCGCTCTCGGACCTTACCTGGTCACGAGGACCGCGATGTCATCC 347
Db 3025 TGCACCTTGGCGCTCTCGGACCTTACCTGGTCACGAGGACCGCGATGTCATCC 3080

RESULT 34
US-08-444-818-137
; Sequence 34 Application US/08444818
; Patent No. 6150087
; GENERAL INFORMATION:
; APPLICANT: Chien, David Y.
; APPLICANT: Rutter, William J.
; TITLE OF INVENTION: NANBV Diagnostics and Vaccines
; NUMBER OF SEQUENCES: 777
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Chiron Corporation
; STREET: 4560 Horton Street
; CITY: Emeryville
; STATE: CA
; COUNTRY: USA
; ZIP: 94608-2916
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/444,818
; FILING DATE:
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/08/403,590
; FILING DATE: 14-MAR-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Harbin, Alisa A.
; REGISTRATION NUMBER: 33,895
; REFERENCE/DOCKET NUMBER: 0110.002
; TELEPHONE: (508)359-3876
; TELEFAX: (508)359-3885

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137

88

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; INFORMATION FOR SEQ ID NO: 137:
; SEQUENCE CHARACTERISTICS:
;   LENGTH: 997 base pairs
;   TYPE: nucleic acid
;   STRANDEDNESS: single
;   TOPOLOGY: linear
;   MOLECULE TYPE: CDNA
;   FEATURE:
;   NAME/KEY: CDS
;   LOCATION: 1..8985
; US-08-444-818-137
;
; Query Match      2.7%; Score 56; DB 3; Length 997;
; Best Local Similarity 100.0%; Pred. No. 1e-18;
; Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
;
Qy 292 TGCACCTTGGCGCTCTCTCGACCTTTACCTGTCACGAGGACGCCGATGTCATTC 347
Db 3367 TGCACCTTGGCGCTCTCTCGACCTTTACCTGTCACGAGGACGCCGATGTCATTC 3422

RESULT 35
US-08-444-818-122
; Sequence 122, Application US/08444818
; Patent No. 6150087
; GENERAL INFORMATION:
; APPLICANT: Chien, David Y.
; APPLICANT: Rutter, William J.
; TITLE OF INVENTION: NANEV Diagnostics and Vaccines
; NUMBER OF SEQUENCES: 777
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Chiron Corporation
; STREET: 4560 Horton Street
; CITY: Emeryville
; STATE: CA
; COUNTRY: USA
; ZIP: 94608-2916
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/444,818
; FILING DATE:
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/08/403,590
; FILING DATE: 14-MAR-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Harbin, Alisa A.
; REGISTRATION NUMBER: 33,895
; REFERENCE/DOCKET NUMBER: 0110.002
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (508)359-3876
; TELEFAX: (508)359-3885
; INFORMATION FOR SEQ ID NO: 123:
; SEQUENCE CHARACTERISTICS:
;   LENGTH: 9185 base pairs
;   TYPE: nucleic acid
;   STRANDEDNESS: single
;   TOPOLOGY: linear
;   MOLECULE TYPE: CDNA
;   ANTI-SENSE: YES
; US-08-444-818-123
;
; Query Match      2.7%; Score 56; DB 3; Length 9185;
; Best Local Similarity 100.0%; Pred. No. 1e-18;
; Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
;
Qy 292 TGCACCTTGGCGCTCTCTCGACCTTTACCTGTCACGAGGACGCCGATGTCATTC 347
Db 5500 TGCACCTTGGCGCTCTCTCGACCTTTACCTGTCACGAGGACGCCGATGTCATTC 5445

RESULT 37
US-08-444-818-176
; Sequence 176, Application US/08444818
; Patent No. 6150087
; GENERAL INFORMATION:
; APPLICANT: Chien, David Y.
; APPLICANT: Rutter, William J.
; TITLE OF INVENTION: NANEV Diagnostics and Vaccines
; NUMBER OF SEQUENCES: 777
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Chiron Corporation
; STREET: 4560 Horton Street
; CITY: Emeryville
; STATE: CA
; COUNTRY: USA
; ZIP: 94608-2916
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk

```

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RESULT 36
US-08-444-818-123/c
; Sequence 123, Application US/08444818
; Patent No. 6150087
; GENERAL INFORMATION:
; APPLICANT: Chien, David Y.
; APPLICANT: Rutter, William J.
; TITLE OF INVENTION: NANEV Diagnostics and Vaccines
; NUMBER OF SEQUENCES: 777
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Chiron Corporation
; STREET: 4560 Horton Street
; CITY: Emeryville
; STATE: CA
; COUNTRY: USA
; ZIP: 94608-2916
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/444,818
; FILING DATE:
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/08/403,590
; FILING DATE: 14-MAR-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Harbin, Alisa A.
; REGISTRATION NUMBER: 33,895
; REFERENCE/DOCKET NUMBER: 0110.002
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (508)359-3876
; TELEFAX: (508)359-3885
; INFORMATION FOR SEQ ID NO: 123:
; SEQUENCE CHARACTERISTICS:
;   LENGTH: 9185 base pairs
;   TYPE: nucleic acid
;   STRANDEDNESS: single
;   TOPOLOGY: linear
;   MOLECULE TYPE: CDNA
;   ANTI-SENSE: YES
; US-08-444-818-123
;
; Query Match      2.7%; Score 56; DB 3; Length 9185;
; Best Local Similarity 100.0%; Pred. No. 1e-18;
; Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
;
Qy 292 TGCACCTTGGCGCTCTCTCGACCTTTACCTGTCACGAGGACGCCGATGTCATTC 347
Db 5500 TGCACCTTGGCGCTCTCTCGACCTTTACCTGTCACGAGGACGCCGATGTCATTC 5445

RESULT 37
US-08-444-818-176
; Sequence 176, Application US/08444818
; Patent No. 6150087
; GENERAL INFORMATION:
; APPLICANT: Chien, David Y.
; APPLICANT: Rutter, William J.
; TITLE OF INVENTION: NANEV Diagnostics and Vaccines
; NUMBER OF SEQUENCES: 777
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Chiron Corporation
; STREET: 4560 Horton Street
; CITY: Emeryville
; STATE: CA
; COUNTRY: USA
; ZIP: 94608-2916
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk

```

176

COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/444,818  
FILING DATE: 14-MAR-1995  
CLASSIFICATION: 424  
PRIOR APPLICATION DATA: US/08/403,590  
APPLICATION NUMBER: 33,895  
FILING DATE: 14-MAR-1995  
ATTORNEY/AGENT INFORMATION:  
NAME: Harbin, Alisa A.  
REGISTRATION NUMBER: 33,895  
REFERENCE/DOCKET NUMBER: 0110.002  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (508)359-3876  
TELEFAX: (508)359-3885  
INFORMATION FOR SEQ ID NO: 176:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 9379 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: cDNA  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: 345  
OTHER INFORMATION: /note= "A heterogeneity may exist at this position which is A or G"  
OTHER INFORMATION: at this position which is A or G"  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: 351  
OTHER INFORMATION: /note= "A heterogeneity may exist at this position which is T or C"  
OTHER INFORMATION: at this position which is T or C"  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: 846  
OTHER INFORMATION: /note= "A heterogeneity may exist at this position which is A or G"  
OTHER INFORMATION: at this position which is A or G"  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: 1319  
OTHER INFORMATION: /note= "A heterogeneity may exist at this position which is A or T"  
OTHER INFORMATION: at this position which is A or T"  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: 2126  
OTHER INFORMATION: /note= "A heterogeneity may exist at this position which is A or C"  
OTHER INFORMATION: at this position which is A or C"  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: 3659  
OTHER INFORMATION: /note= "A heterogeneity may exist at this position which is C or T"  
OTHER INFORMATION: at this position which is C or T"  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: 3689  
OTHER INFORMATION: /note= "A heterogeneity may exist at this position which is G or C"  
OTHER INFORMATION: at this position which is G or C"  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: 4146  
OTHER INFORMATION: /note= "A heterogeneity may exist at this position which is C or T"  
OTHER INFORMATION: at this position which is C or T"  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: 4680  
OTHER INFORMATION: /note= "A heterogeneity may exist at this position which is G or A"  
OTHER INFORMATION: at this position which is G or A"  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: 9080  
OTHER INFORMATION: /note= "A heterogeneity may exist at this position which is A or G"  
OTHER INFORMATION: at this position which is A or G"

OTHER INFORMATION: at this position which is A or G"  
US-08-444-818-176  
Query Match 2.7%; Score 56; DB 3; Length 9379;  
Best Local Similarity 100.0%; Pred. No. 1e-18;  
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 292 TGCACCTTCGGCTCCTCGGACCTTTACCTGCTCAGGACGACGCGGATGTCATTC 347  
DB 3686 TGCACCTTCGGCTCCTCGGACCTTTACCTGCTCAGGACGACGCGGATGTCATTC 3741  
RESULT 38  
US-09-388-874-1  
Sequenced Application US/093888974  
Patent No. 6284249  
GENERAL INFORMATION:  
APPLICANT: Veronique Barban  
TITLE OF INVENTION: VACCINE COMPOSITION FOR PREVENTING OR  
FILE REFERENCE: PMCF97-03A  
CURRENT APPLICATION NUMBER: US/09/388,874  
CURRENT FILING DATE: 1999-09-02  
EARLIER APPLICATION NUMBER: PCT/FR98/00448  
EARLIER FILING DATE: 1998-03-06  
EARLIER APPLICATION NUMBER: 97/02,887  
EARLIER FILING DATE: 1997-03-06  
NUMBER OF SEQ ID NOS: 2  
SOFTWARE: FastSeq for Windows Version 3.0  
SEQ ID NO 1  
LENGTH: 9379  
TYPE: DNA  
ORGANISM: Virus  
FEATURE:  
NAME/KEY: CDS  
LOCATION: (320)...(9352)  
US-09-388-874-1

Query Match 2.7%; Score 56; DB 3; Length 9379;  
Best Local Similarity 100.0%; Pred. No. 1e-18;  
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 292 TGCACCTTCGGCTCCTCGGACCTTTACCTGCTCAGGACGACGCGGATGTCATTC 347  
DB 3686 TGCACCTTCGGCTCCTCGGACCTTTACCTGCTCAGGACGACGCGGATGTCATTC 3741

RESULT 39  
US-09-916-359-1  
Sequence 1, Application US/09916359  
Patent No. 6538123  
GENERAL INFORMATION:  
APPLICANT: Veronique Barban  
TITLE OF INVENTION: VACCINE COMPOSITION FOR PREVENTING OR  
FILE REFERENCE: PMCF97-03A  
CURRENT APPLICATION NUMBER: US/09/916,359  
CURRENT FILING DATE: 2001-07-26  
PRIOR APPLICATION NUMBER: 09/388,874  
PRIOR FILING DATE: 1999-09-02  
PRIOR APPLICATION NUMBER: 97/02,887  
PRIOR FILING DATE: 1997-03-06  
NUMBER OF SEQ ID NOS: 2  
SOFTWARE: FastSeq for Windows Version 3.0  
SEQ ID NO 1  
LENGTH: 9379  
TYPE: DNA  
ORGANISM: Virus  
FEATURE:  
NAME/KEY: CDS  
LOCATION: (320)...(9352)  
US-09-916-359-1

whole virus 102 (2)  
sep 2001

Gene

Query Match 2.7%; Score 56; DB 4; Length 9379;  
 Best Local Similarity 100.0%; Pred. No. 1e-18; Indels 0; Gaps 0;  
 Matches 56; Conservative 0; Mismatches 0;  
 QY 292 TGCACCTTCGGCTCTCTGGACCTTACTGTGTCACGAGGACGCGGATGTCATCC 347  
 Db 3686 TGCACCTTCGGCTCTCTGGACCTTACTGTGTCACGAGGACGCGGATGTCATCC 3741

RESULT 40  
 US-07-910-760-9  
 ; Sequence(s) Application US/07910760  
 ; Patent No. 5683864  
 ; GENERAL INFORMATION:  
 ; APPLICANT: Houghton, Michael  
 ; APPLICANT: Choo, Qui-Lim  
 ; APPLICANT: Kuo, George  
 ; TITLE OF INVENTION: Combinations of Hepatitis C virus (HCV)  
 ; TITLE OF INVENTION: Antigens for Use in Immunoassays for Anti-HCV Antibodies  
 ; NUMBER OF SEQUENCES: 12  
 ; CORRESPONDENCE ADDRESS:  
 ; ADDRESSEE: Chiron Corporation  
 ; STREET: P.O. Box 8097 (Int. Prop. R-440)  
 ; CITY: Emeryville  
 ; STATE: CA  
 ; COUNTRY: U.S.A.  
 ; ZIP: 94662-8097  
 ; COMPUTER READABLE FORM:  
 ; MEDIUM TYPE: Floppy disk  
 ; COMPUTER: IBM PC compatible  
 ; OPERATING SYSTEM: PC-DOS/MS-DOS  
 ; SOFTWARE: Patent in Release #1.0, Version #1.25  
 ; CURRENT APPLICATION DATA:  
 ; APPLICATION NUMBER: US/07910,760  
 ; FILING DATE: 07-JUL-1992  
 ; CLASSIFICATION: 435  
 ; ATTORNEY/AGENT INFORMATION:  
 ; NAME: Blackburn Esq., Robert P.  
 ; REGISTRATION NUMBER: 30,447  
 ; REFERENCE/DOCKET NUMBER: 0101.002  
 ; TELECOMMUNICATION INFORMATION:  
 ; TELEPHONE: (510) 601-2702  
 ; TELEFAX: (510) 655-3542  
 ; INFORMATION FOR SEQ ID NO: 9:  
 ; SEQUENCE CHARACTERISTICS:  
 ; LENGTH: 9401 base pairs  
 ; TYPE: nucleic acid  
 ; STRANDEDNESS: double  
 ; TOPOLOGY: linear  
 ; MOLECULE TYPE: DNA (genomic)  
 ; FEATURE:  
 ; NAME/KEY: CDS  
 ; LOCATION: 342..9374  
 ; FEATURE:  
 ; NAME/KEY: misc\_feature  
 ; LOCATION: 366  
 ; OTHER INFORMATION: /note= "This amino acid position  
 ; OTHER INFORMATION: can also be Arg."  
 ; FEATURE:  
 ; NAME/KEY: misc\_feature  
 ; LOCATION: 372  
 ; OTHER INFORMATION: /note= "This amino acid position/  
 ; OTHER INFORMATION: can also be Thr."  
 ; FEATURE:  
 ; NAME/KEY: misc\_feature  
 ; LOCATION: 867  
 ; OTHER INFORMATION: /note= "This amino acid position  
 ; OTHER INFORMATION: can also be Thr."  
 ; FEATURE:  
 ; NAME/KEY: misc\_feature  
 ; LOCATION: 1341  
 ; OTHER INFORMATION: /note= "This amino acid position  
 ; OTHER INFORMATION: can also be Val."

FEATURE:  
 NAME/KEY: misc\_feature  
 LOCATION: 2148  
 OTHER INFORMATION: /note= "This amino acid position  
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 FEATURE:  
 NAME/KEY: misc\_feature  
 LOCATION: 2883  
 OTHER INFORMATION: /note= "This amino acid position  
 OTHER INFORMATION: can also be Asn."  
 FEATURE:  
 NAME/KEY: misc\_feature  
 LOCATION: 3681  
 OTHER INFORMATION: /note= "This amino acid position  
 OTHER INFORMATION: can also be Ser."  
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 NAME/KEY: misc\_feature  
 LOCATION: 3690  
 OTHER INFORMATION: /note= "This amino acid position  
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 NAME/KEY: misc\_feature  
 LOCATION: 4167  
 OTHER INFORMATION: /note= "This amino acid position  
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 NAME/KEY: misc\_feature  
 LOCATION: 4323  
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 NAME/KEY: misc\_feature  
 LOCATION: 4701  
 OTHER INFORMATION: /note= "This amino acid position  
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 FEATURE:  
 NAME/KEY: misc\_feature  
 LOCATION: 4752  
 OTHER INFORMATION: /note= "This amino acid position  
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 FEATURE:  
 NAME/KEY: misc\_feature  
 LOCATION: 5970  
 OTHER INFORMATION: /note= "This amino acid position  
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 NAME/KEY: misc\_feature  
 LOCATION: 6183  
 OTHER INFORMATION: /note= "This amino acid position  
 OTHER INFORMATION: can also be His."  
 FEATURE:  
 NAME/KEY: misc\_feature  
 LOCATION: 6186  
 OTHER INFORMATION: /note= "This amino acid position  
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 FEATURE:  
 NAME/KEY: misc\_feature  
 LOCATION: 6402  
 OTHER INFORMATION: /note= "This amino acid position  
 OTHER INFORMATION: can also be Val."  
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 NAME/KEY: misc\_feature  
 LOCATION: 7386  
 OTHER INFORMATION: /note= "This amino acid position  
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 FEATURE:  
 NAME/KEY: misc\_feature  
 LOCATION: 7494  
 OTHER INFORMATION: /note= "This amino acid position  
 OTHER INFORMATION: can also be Phe."  
 FEATURE:  
 NAME/KEY: misc\_feature  
 LOCATION: 7497

*the virus? whole virus*

3768  
3705

OTHER INFORMATION: /note= "This amino acid position  
OTHER INFORMATION: can also be Ala."  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: 7845  
OTHER INFORMATION: /note= "This amino acid position  
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LOCATION: 8409  
OTHER INFORMATION: /note= "This amino acid position  
OTHER INFORMATION: can also be Gly."  
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LOCATION: 9102  
OTHER INFORMATION: /note= "This amino acid position  
OTHER INFORMATION: can also be Gly."  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: 9327  
OTHER INFORMATION: /note= "This amino acid position  
OTHER INFORMATION: can also be Pro."  
US-07-910-760-9

Query Match 2.7% Score 56; DB 1; Length 9401;  
Best Local Similarity 100.0%; Pred. No. 18-18;  
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0  
QY 292 TGACATTCGGCGCTCTCGGACCTTTACCTGGTCACGAGGACCGCGATGTCATTCC 347  
Db 3708 TGACATTCGGCGCTCTCGGACCTTTACCTGGTCACGAGGACCGCGATGTCATTCC 3763

*Save as before*

RESULT 41  
US-08-440-549-9  
Sequence 9 Application US/08440519  
Patent No 5712087  
GENERAL INFORMATION:  
APPLICANT: Houghton, Michael  
APPLICANT: Choo, Qui-Lim  
APPLICANT: Kuo, George  
TITLE OF INVENTION: Combinations of Hepatitis C virus (HCV)  
TITLE OF INVENTION: Antigens for Use in Immunoassays for Anti-HCV Antibodies  
NUMBER OF SEQUENCES: 12  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Chiron Corporation  
STREET: P.O. Box 8097 (Int. Prop. R-440)  
CITY: Emeryville  
STATE: CA  
COUNTRY: U.S.A.  
ZIP: 94662-8097  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patent In Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/440,519  
FILING DATE: 12-MAY-1995  
CLASSIFICATION: 530  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/910,760  
FILING DATE: 07-JUL-1992  
ATTORNEY/AGENT INFORMATION:  
NAME: Blackburn Esq., Robert P.  
REGISTRATION NUMBER: 30,447  
REFERENCE/DOCKET NUMBER: 0101.002  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (510) 601-2702  
TELEFAX: (510) 655-3542  
INFORMATION FOR SEQ ID NO: 9:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 9401 base pairs

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; NAME/KEY: misc_feature
; LOCATION: 6183
; OTHER INFORMATION: /note= "This amino acid position
; OTHER INFORMATION: can also be His."
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 6186
; OTHER INFORMATION: /note= "This amino acid position
; OTHER INFORMATION: can also be Cys."
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 6402
; OTHER INFORMATION: /note= "This amino acid position
; OTHER INFORMATION: can also be Val."
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 7386
; OTHER INFORMATION: /note= "This amino acid position
; OTHER INFORMATION: can also be Ser."
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 7494
; OTHER INFORMATION: /note= "This amino acid position
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; NAME/KEY: misc_feature
; LOCATION: 7497
; OTHER INFORMATION: /note= "This amino acid position
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; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 7845
; OTHER INFORMATION: /note= "This amino acid position
; OTHER INFORMATION: can also be Phe."
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 8409
; OTHER INFORMATION: /note= "This amino acid position
; OTHER INFORMATION: can also be Gly."
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 9102
; OTHER INFORMATION: /note= "This amino acid position
; OTHER INFORMATION: can also be Gly."
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 9327
; OTHER INFORMATION: /note= "This amino acid position
; OTHER INFORMATION: can also be Pro."

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US-08-440-519-9

Query Match 2.7%; Score 56; DB 1; Length 9401;  
 Best Local Similarity 100.0%; Pred. No. 1e-18;  
 Matches 56; Conservative 0; Mismatches 0; Gaps 0;

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Qy 292 TGCACCTGGCGCTCCTCGGACCTTACCTGTCACGAGCAGCCGATGTCATCC 347
Db 3708 TGCACCTGGCGCTCCTCGGACCTTACCTGTCACGAGCAGCCGATGTCATCC 3763

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RESULT 42

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US-08-440-519-9
; Sequence 9, Application US/08440549
; Patent No. 6312889
; GENERAL INFORMATION:
; APPLICANT: Houghton, Michael
; APPLICANT: Choo, Qui-Lim
; APPLICANT: Kuo, George
; TITLE OF INVENTION: Combinations of Hepatitis C virus (HCV)
; TITLE OF INVENTION: Antigens for Use in Immunoassays for Anti-HCV Antibodies
; NUMBER OF SEQUENCES: 12
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Chiron Corporation

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; STREET: P.O. Box 8097 (Int. Prop. R-440)
; CITY: Emeryville
; STATE: CA
; COUNTRY: U.S.A.
; ZIP: 94662-8097
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/440,549
; FILING DATE: 12-MAY-1995
; CLASSIFICATION: 435
; PRIORITY APPLICATION DATA:
; APPLICATION NUMBER: US 07/910,760
; FILING DATE: 07-JUL-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Blackburn Esq., Robert P.
; REGISTRATION NUMBER: 30,447
; REFERENCE/DOCKET NUMBER: 0101.002
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (510) 601-2702
; TELEFAX: (510) 635-3542
; INFORMATION FOR SEQ ID NO: 9:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 9401 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 342..9374
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 366
; OTHER INFORMATION: /note= "This amino acid position
; OTHER INFORMATION: can also be Arg."
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 372
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; NAME/KEY: misc_feature
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; OTHER INFORMATION: /note= "This amino acid position
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; LOCATION: 2148
; OTHER INFORMATION: /note= "This amino acid position
; OTHER INFORMATION: can also be Ile."
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 2883
; OTHER INFORMATION: /note= "This amino acid position
; OTHER INFORMATION: can also be Asn."
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 3681
; OTHER INFORMATION: /note= "This amino acid position
; OTHER INFORMATION: can also be Ser."
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 3690
; OTHER INFORMATION: /note= "This amino acid position

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OTHER INFORMATION: can also be Thr."  
NAME/KEY: misc\_feature  
LOCATION: 4167  
OTHER INFORMATION: /note= "This amino acid position  
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NAME/KEY: misc\_feature  
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NAME/KEY: misc\_feature  
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FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: 6402  
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OTHER INFORMATION: can also be Ser."  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: 7494  
OTHER INFORMATION: /note= "This amino acid position  
OTHER INFORMATION: can also be Phe."  
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NAME/KEY: misc\_feature  
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OTHER INFORMATION: /note= "This amino acid position  
OTHER INFORMATION: can also be Phe."  
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NAME/KEY: misc\_feature  
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NAME/KEY: misc\_feature  
LOCATION: 9102  
OTHER INFORMATION: /note= "This amino acid position  
OTHER INFORMATION: can also by Gly."  
FEATURE:  
NAME/KEY: misc\_feature

LOCATION: 9327  
OTHER INFORMATION: /note= "This amino acid position  
OTHER INFORMATION: can also be Pro."  
US-08-440-549-9  
Query Match 2.7%; Score 56; DB 4; Length 9401;  
Best Local Similarity 100.0%; Pred. No. 1e-18;  
Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 292 TGCACCTTGGGCTCTCTCGGACCTTTACCTGTGTCACGAGGACGCGGATGTCATTCC 347  
DB 3708 TGCACCTTGGGCTCTCTCGGACCTTTACCTGTGTCACGAGGACGCGGATGTCATTCC 3763

RESULT 43  
US-08-823-895A-25  
Sequence 25 Application US/08823895A  
Patent No. 6433159  
GENERAL INFORMATION:  
APPLICANT: Kevin P. Anderson  
TITLE OF INVENTION: Compositions And Methods For  
TREATMENT OF HEPATITIS C VIRUS-ASSOCIATED DISEASES  
NUMBER OF SEQUENCES: 27  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Jane Massey Licata, Esq.  
STREET: 66 E. Main Street  
CITY: Marlton  
STATE: NJ  
COUNTRY: USA  
ZIP: 08053  
COMPUTER READABLE FORM:  
MEDIUM TYPE: DISKETTE, 3.5 INCH, 1.44 MB STORAGE  
COMPUTER: IBM 486  
OPERATING SYSTEM: WINDOWS FOR WORKGROUPS  
SOFTWARE: WORDPERFECT 5.1  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/823.895A  
FILING DATE: March 17, 1997  
CLASSIFICATION: 514  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/453,085  
FILING DATE: May 30, 1995  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 07/945,289  
FILING DATE: September 10, 1992  
ATTORNEY/AGENT INFORMATION:  
NAME: Jane Massey Licata  
REGISTRATION NUMBER: 32,257  
REFERENCE/DOCKET NUMBER: ISPH-0203  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (609) 779-2400  
TELEFAX: (609) 810-1454  
INFORMATION FOR SEQ ID NO: 25:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 9401  
TYPE: Nucleic  
STRANDEDNESS: Single  
TOPOLOGY: Linear  
ANTI-SENSE: NO  
US-08-823-895A-25

Query Match 2.7%; Score 56; DB 4; Length 9401;  
Best Local Similarity 100.0%; Pred. No. 1e-18;  
Matches 56; Conservative 0; Mismatches 0; Indels 0;  
QY 292 TGCACCTTGGGCTCTCTCGGACCTTTACCTGTGTCACGAGGACGCGGATGTCATTCC 347  
DB 3708 TGCACCTTGGGCTCTCTCGGACCTTTACCTGTGTCACGAGGACGCGGATGTCATTCC 3763

RESULT 44  
PCT-US91-02225-9  
Sequence 9 Application PC/TUS9102225

10225-9  
who's virus  
15/10/97

GENERAL INFORMATION:  
 APPLICANT: HOUGHTON, MICHAEL  
 APPLICANT: CHOO, QUI-LIM  
 APPLICANT: KOO, GEORGE  
 TITLE OF INVENTION: COMBINATIONS OF HEPATITIS C VIRUS  
 TITLE OF INVENTION: ANTIBODIES  
 NUMBER OF SEQUENCES: 10  
 CORRESPONDENCE ADDRESS:  
 ADDRESSEE: Morrison & Foerster  
 STREET: 545 Middlefield Road, Suite 200  
 CITY: Menlo Park  
 STATE: CA  
 COUNTRY: USA  
 ZIP: 94025  
 COMPUTER READABLE FORM:  
 MEDIUM TYPE: Floppy disk  
 COMPUTER: IBM PC compatible  
 OPERATING SYSTEM: PC-DOS/MS-DOS  
 SOFTWARE: PatentIn Release #1.0, Version #1.25  
 CURRENT APPLICATION DATA:  
 APPLICATION NUMBER: PCT/US91/02225  
 FILING DATE: 19910329  
 CLASSIFICATION: 435  
 ATTORNEY/AGENT INFORMATION:  
 NAME: CIOTTI, THOMAS E.  
 REGISTRATION NUMBER: 21,013  
 REFERENCE/DOCKET NUMBER: 2300-0101.44  
 TELECOMMUNICATION INFORMATION:  
 TELEPHONE: (415) 327-7250  
 TELEFAX: (415) 327-2951  
 TELEX: 706141  
 INFORMATION FOR SEQ ID NO: 9:  
 SEQUENCE CHARACTERISTICS:  
 LENGTH: 9401 base pairs  
 TYPE: NUCLEIC ACID  
 STRANDEDNESS: unknown  
 TOPOLOGY: unknown  
 MOLECULE TYPE: DNA (genomic)  
 PCT-US91-02225-9

Query Match 2.7%; Score 56; DB 5; Length 9401;  
 Best Local Similarity 100.0%; Pred. No. 1e-18;  
 Matches 56; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 292 TGCACTGGCGCTCCTCGGACCTTACTGCTACGAGGACCGCGATGTCATCC 347  
 Db 3708 TGCACTGGCGCTCCTCGGACCTTACTGCTACGAGGACCGCGATGTCATCC 3763

RESULT 45  
 US-08-823-895A-26  
 Sequence 26, Application US/08823895A  
 Patent No. 6433159  
 GENERAL INFORMATION:  
 APPLICANT: Kevin P. Anderson  
 TITLE OF INVENTION: Compositions And Methods For  
 TITLE OF INVENTION: Treatment Of Hepatitis C Virus-Associated Diseases  
 NUMBER OF SEQUENCES: 27  
 CORRESPONDENCE ADDRESS:  
 ADDRESSEE: Jane Massey Licata, Esq.  
 STREET: 66 E. Main Street  
 CITY: Marlton  
 STATE: NJ  
 COUNTRY: USA  
 ZIP: 08053  
 COMPUTER READABLE FORM:  
 MEDIUM TYPE: DISKETTE, 3.5 INCH, 1.44 Mb STORAGE  
 COMPUTER: IBM 486  
 OPERATING SYSTEM: WINDOWS FOR WORKGROUPS  
 SOFTWARE: WORDPERFECT 5.1  
 CURRENT APPLICATION DATA:  
 APPLICATION NUMBER: US/08/823,895A

FILING DATE: March 17, 1997  
 CLASSIFICATION: 514  
 PRIOR APPLICATION DATA:  
 APPLICATION NUMBER: 08/453,085  
 FILING DATE: May 30, 1995  
 PRIOR APPLICATION DATA:  
 APPLICATION NUMBER: 07/945,289  
 FILING DATE: September 10, 1992  
 ATTORNEY/AGENT INFORMATION:  
 NAME: Jane Massey Licata  
 REGISTRATION NUMBER: 32,257  
 REFERENCE/DOCKET NUMBER: ISPH-0203  
 TELECOMMUNICATION INFORMATION:  
 TELEPHONE: (609) 779-2400  
 TELEFAX: (609) 810-1454  
 INFORMATION FOR SEQ ID NO: 26:  
 SEQUENCE CHARACTERISTICS:  
 LENGTH: 9416  
 TYPE: Nucleic  
 STRANDEDNESS: Single  
 TOPOLOGY: Linear  
 ANTI-SENSE: NO  
 US-08-823-895A-26

Query Match 2.6%; Score 53; DB 4; Length 9416;  
 Best Local Similarity 100.0%; Pred. No. 3.5e-17;  
 Matches 53; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 598 GTGGCCACCTGTCATGCTCCACCGGACGCGTAAGACCAAGGTCCTCCGC 650  
 Db 4014 GTGGCCACCTGTCATGCTCCACCGGACGCGTAAGACCAAGGTCCTCCGC 4066

RESULT 46  
 US-10-104-966-13  
 Sequence 13, Application US/10104966  
 Patent No. 6680054  
 GENERAL INFORMATION:  
 APPLICANT: Matti Sallberg  
 APPLICANT: Catharina Hultgren  
 TITLE OF INVENTION: VACCINES CONTAINING RIBAVIRIN AND  
 TITLE OF INVENTION: METHODS OF USE THEREOF  
 FILE REFERENCE: TRIPEP.23AUSCI  
 CURRENT APPLICATION NUMBER: US/10/104,966  
 CURRENT FILING DATE: 2002-03-22  
 PRIOR APPLICATION NUMBER: 09/705,547  
 PRIOR FILING DATE: 2000-11-03  
 PRIOR APPLICATION NUMBER: 60/229,175  
 PRIOR FILING DATE: 2000-08-29  
 NUMBER OF SEQ ID NOS: 15  
 SOFTWARE: FastSeq for Windows Version 4.0  
 SEQ ID NO 13  
 LENGTH: 9416  
 TYPE: DNA  
 ORGANISM: Artificial Sequence  
 FEATURE:  
 OTHER INFORMATION: Hepatitis C virus sequence  
 US-10-104-966-13

Query Match 2.6%; Score 53; DB 4; Length 9416;  
 Best Local Similarity 100.0%; Pred. No. 3.5e-17;  
 Matches 53; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 598 GTGGCCACCTGTCATGCTCCACCGGACGCGTAAGACCAAGGTCCTCCGC 650  
 Db 4014 GTGGCCACCTGTCATGCTCCACCGGACGCGTAAGACCAAGGTCCTCCGC 4066

RESULT 47  
 US-09-014-416-2  
 Sequence 2, Application US/09014416  
 Patent No. 6153421  
 GENERAL INFORMATION:

13

iso lat

4066  
4066  
4066

APPLICANT: Yanagi, Masayuki  
APPLICANT: Bukh, Jens  
APPLICANT: Emerson, Susanne U.  
APPLICANT: Purcell, Robert H.  
TITLE OF INVENTION: CLONED GENOMES OF INFECTIOUS HEPATITIS C VIRUSES AND  
TITLE OF INVENTION: USES THEREOF  
FILE REFERENCE: 20264276  
CURRENT APPLICATION NUMBER: US/09/014,416  
CURRENT FILING DATE: 1998-01-27  
EARLIER APPLICATION NUMBER: US 60/053,062  
EARLIER FILING DATE: 1997-07-18  
NUMBER OF SEQ ID NOS: 65  
SOFTWARE: PatentIn Ver. 2.1  
SEQ ID NO 2  
LENGTH: 9599  
TYPE: DNA  
ORGANISM: Hepatitis C virus  
US-09-014-416-2

Query Match 2.6%; Score 53; DB 3; Length 9599;  
Best Local Similarity 100.0%; Pred. No. 3.5e-17; Indels 0; Gaps 0;  
Matches 53; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 598 GTGGCCCACTGCTGCTCCACCGGCGGTAAGAGCACCAGGTCCTCCGGC 650  
Db 4014 GTGGCCCACTGCTGCTCCACCGGCGGTAAGAGCACCAGGTCCTCCGGC 4066

RESULT 48  
US-09-014-416-6  
Sequence 6, Application US/09014416  
Patent No. 6153421  
GENERAL INFORMATION:  
APPLICANT: Yanagi, Masayuki  
APPLICANT: Bukh, Jens  
APPLICANT: Emerson, Susanne U.  
APPLICANT: Purcell, Robert H.  
TITLE OF INVENTION: CLONED GENOMES OF INFECTIOUS HEPATITIS C VIRUSES AND  
TITLE OF INVENTION: USES THEREOF  
FILE REFERENCE: 20264276  
CURRENT APPLICATION NUMBER: US/09/014,416  
CURRENT FILING DATE: 1998-01-27  
EARLIER APPLICATION NUMBER: US 60/053,062  
EARLIER FILING DATE: 1997-07-18  
NUMBER OF SEQ ID NOS: 65  
SOFTWARE: PatentIn Ver. 2.1  
SEQ ID NO 6  
LENGTH: 9599  
TYPE: DNA  
ORGANISM: Hepatitis C virus  
US-09-014-416-6

Query Match 2.6%; Score 53; DB 3; Length 9599;  
Best Local Similarity 100.0%; Pred. No. 3.5e-17; Indels 0; Gaps 0;  
Matches 53; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 598 GTGGCCCACTGCTGCTCCACCGGCGGTAAGAGCACCAGGTCCTCCGGC 650  
Db 4014 GTGGCCCACTGCTGCTCCACCGGCGGTAAGAGCACCAGGTCCTCCGGC 4066

RESULT 49  
US-09-014-416-1  
Sequence 1, Application US/08811566  
Patent No. 6127116  
GENERAL INFORMATION:  
APPLICANT: Rice, Charles et al.  
TITLE OF INVENTION: FUNCTIONAL DNA CLONE FOR HEPATITIS C  
TITLE OF INVENTION: VIRUS (HCV) AND USES THEREOF  
NUMBER OF SEQUENCES: 21  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: David A. Jackson, Esq.  
STREET: 411 Hackensack Ave, Continental Plaza, 4th

STREET: Floor  
CITY: Hackensack  
STATE: New Jersey  
COUNTRY: USA  
ZIP: 07601  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.30  
CURRENT APPLICATION DATA: US/08/811,566  
APPLICATION NUMBER: US/08/811,566  
FILING DATE: 03-MAR-1997  
CLASSIFICATION: 435  
ATTORNEY/AGENT INFORMATION:  
NAME: Jackson Esq., David A.  
REGISTRATION NUMBER: 26,742  
REFERENCE/DOCKET NUMBER: 1113-1-006  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 201-487-5800  
TELEFAX: 201-343-1684  
INFORMATION FOR SEQ ID NO: 1:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 9646 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: cDNA  
HYPOTHETICAL: NO  
ANTI-SENSE: NO  
US-08-811-566-1

Query Match 2.6%; Score 53; DB 3; Length 9646;  
Best Local Similarity 100.0%; Pred. No. 3.5e-17; Indels 0; Gaps 0;  
Matches 53; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 598 GTGGCCCACTGCTGCTCCACCGGCGGTAAGAGCACCAGGTCCTCCGGC 650  
Db 4014 GTGGCCCACTGCTGCTCCACCGGCGGTAAGAGCACCAGGTCCTCCGGC 4066

RESULT 50  
US-09-034-756-1  
Sequence 1, Application US/09034756  
Patent No. 6332028  
GENERAL INFORMATION:  
APPLICANT: Rice, Charles et al.  
TITLE OF INVENTION: FUNCTIONAL DNA CLONE FOR HEPATITIS C  
TITLE OF INVENTION: VIRUS (HCV) AND USES THEREOF  
NUMBER OF SEQUENCES: 21  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: HOWELL & HAFFKAMP, L.C.  
STREET: 7733 FORSYTH BLVD., SUITE 1400  
CITY: ST. LOUIS  
STATE: MO  
COUNTRY: USA  
ZIP: 63105  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/034,756  
FILING DATE: 04-May-1998  
CLASSIFICATION: <Unknown>  
ATTORNEY/AGENT INFORMATION:  
NAME: HOLLAND, DONALD R.  
REGISTRATION NUMBER: 35,197  
REFERENCE/DOCKET NUMBER: 6029-4831  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 314-727-5188  
TELEFAX: 314-727-6092

4066  
4066

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; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
;   LENGTH: 9646 base pairs
;   TYPE: nucleic acid
;   STRANDEDNESS: double
;   TOPOLOGY: linear
; MOLECULE TYPE: cDNA
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; SEQUENCE DESCRIPTION: SEQ ID NO: 1:
US-09-034-756-1

Query Match      2.6%; Score 53; DB 4; Length 9646;
Best Local Similarity 100.0%; Pred. No. 3.5e-17;
Matches 53; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 598 GTGGCCCACTGTCATGCTCCACCGGCGAGGTAAGAGCACCACCAAGGTCCCGGC 650
    |||||||
Db 4014 GTGGCCCACTGTCATGCTCCACCGGCGAGGTAAGAGCACCACCAAGGTCCCGGC 4066
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Search completed: August 19, 2004, 10:24:16  
Job time : 177 secs

Search completed: August 19, 2004, 10:14:09  
Job time : 4815 secs

GenCore version 5.1.6  
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OM nucleic - nucleic search, using sw model

Run on: August 19, 2004, 04:53:05 ; Search time 4815 Seconds  
(without alignments)  
12782.130 Million cell updates/sec

Title: US-09-930-591-1  
Perfect score: 2061  
Sequence: 1 atggcgctatcacggccta.....atgaatggaagagtgtga 2061

Scoring table: OLIGO\_NUC  
Gapop 60.0 , Gapext 60.0

Searched: 27513289 seqs, 14931090276 residues

Word size : 35

Total number of hits satisfying chosen parameters: 0

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Listing first 1000 summaries

Database : EST.\*  
1: em\_estba:\*  
2: em\_esthum:\*  
3: em\_estin:\*  
4: em\_estmu:\*  
5: em\_estov:\*  
6: em\_estpl:\*  
7: em\_estro:\*  
8: em\_htc:\*  
9: gb\_est1:\*  
10: gb\_est2:\*  
11: gb\_htc:\*  
12: gb\_est3:\*  
13: gb\_est4:\*  
14: gb\_est5:\*  
15: em\_estfun:\*  
16: em\_estom:\*  
17: em\_gss\_hum:\*  
18: em\_gss\_inv:\*  
19: em\_gss\_pln:\*  
20: em\_gss\_vrt:\*  
21: em\_gss\_fun:\*  
22: em\_gss\_mam:\*  
23: em\_gss\_mus:\*  
24: em\_gss\_pro:\*  
25: em\_gss\_rod:\*  
26: em\_gss\_phg:\*  
27: em\_gss\_vrt:\*  
28: gb\_gss1:\*  
29: gb\_gss2:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
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No matches found

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